

PCT

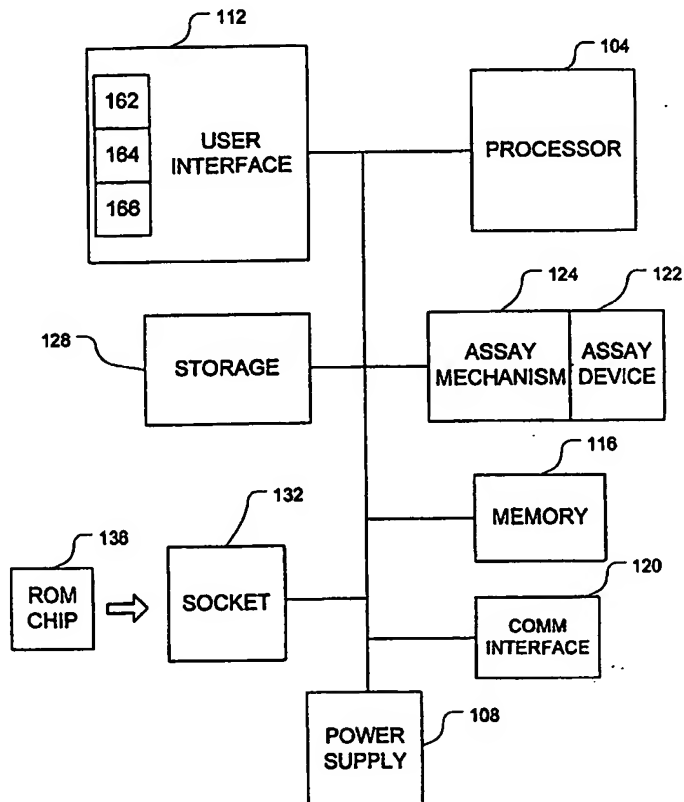
WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification G01N 21/64	A1	(11) International Publication Number: WO 99/35486 (43) International Publication Date: 15 July 1999 (15.07.99)
(21) International Application Number: PCT/US98/26576 (22) International Filing Date: 15 December 1998 (15.12.98) (30) Priority Data: 09/003,090 5 January 1998 (05.01.98) US (63) Related by Continuation (CON) or Continuation-in-Part (CIP) to Earlier Application US 09/003,090 (CIP) Filed on 5 January 1998 (05.01.98) (71) Applicant (for all designated States except US): BIOSITE DIAGNOSTICS, INC. [US/US]; Suite D, 11030 Roselle Street, San Diego, CA 92121 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): BUECHLER, Kenneth, F. [US/US]; 12523 Manifesto Place, San Diego, CA 92130 (US). ANDERBERG, Joseph, M. [US/US]; 470 Delage Court, Encinitas, CA 92024 (US). MCPHERSON, Paul, H. [US/US]; 1449 Elva Court, Encinitas, CA 92024 (US).		(74) Agents: YANNUZZI, Daniel, N. et al.; Lyon & Lyon LLP, Suite 4700, 633 West Fifth Street, Los Angeles, CA 90071-2066 (US). (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>

(54) Title: IMMUNOASSAY FLUOROMETER**(57) Abstract**

A fluorometer for sensing the fluorescence of a sample utilizes an optical energy source for exciting a sample to be tested and an optical energy detector for detecting the emitted energy from the excited sample. Drive electronics are used for positioning the sample with respect to the optical components allowing a plurality of sample regions to be tested. A processor is utilized to control the operation of the test in accordance with test instructions and for processing the emitted energy detected from the sample to determine test results. A ROM chip socket accepts a plurality of ROM chips, wherein each ROM chip stores test data sets for one or more test types to be performed. ROM chips can be swapped to allow the fluorometer to be configured and reconfigured to perform a plurality of different tests. A communications interface facilitates the sharing of test information between the fluorometer and external entities.



FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TC	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon	KR	Republic of Korea	PL	Poland		
CN	China	KZ	Kazakhstan	PT	Portugal		
CU	Cuba	LC	Saint Lucia	RO	Romania		
CZ	Czech Republic	LI	Liechtenstein	RU	Russian Federation		
DE	Germany	LK	Sri Lanka	SD	Sudan		
DK	Denmark	LR	Liberia	SE	Sweden		
EE	Estonia			SG	Singapore		

DESCRIPTIONIMMUNOASSAY FLUOROMETER5 RELATED APPLICATIONS

The present application is related to copending United States Patent Applications No. 09/003,066, docket number 230/004, titled "Media Carrier for an Assay Device," and No. To Be Assigned, docket number 230/006, titled "Methods for Monitoring the Status of Assays and Immunoassays," each of which are filed concurrently herewith and each of
10 which are incorporated herein by reference in their entirety.

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates generally to fluorometers, and more particularly to a
15 system and method for performing automated fluorescent measurements.

2. Related Art

Conventional medical devices used to perform fluorescent readings are large, dedicated machines. Typical fluorometers are bench-top devices which are not easily transported from location to location. Additionally, conventional fluorometers are not
20 easily capable of being programmed by the user or otherwise configured by the user to perform a plurality of different types of tests. Instead, the conventional fluorometer is factory-programmed to perform a predefined test protocol.

To perform a test using a conventional fluorometer, a laboratory technician obtains a sample. The sample can be a biological fluid, such as blood, serum, plasma, urine, a
25 fecal extract and the like or it can be an environmental sample, such as water, a ground extract, a chemical and the like or it can be an extract of a food product. In the case of a blood sample, the blood is first separated into plasma or serum, which becomes the sample, and cellular fractions using a centrifuge. The sample is then generally deposited into a small test tube that is inserted into the fluorometer. Because the conventional
30 fluorometer can accept several samples, the technician enters an identification of the sample and the location of the test tube into the fluorometer.

Once one or more test tubes containing samples are positioned in the fluorometer, the test begins. Contemporary fluorometers use robotics to pipet the sample and the

reagents and to position one or more optical sensors by the sample to obtain the necessary readings. The readings are recorded along with the associated test-tube location designations. The location designation is used to identify the sample.

5 SUMMARY OF THE INVENTION

The present invention provides a system and method for performing automated fluorometry. According to the invention, a fluorometer is provided which includes functionality to provide enhanced operational characteristics for the measurement of analytes in a sample. The system and method has particular importance as a laboratory or
10 non-laboratory tool for rapidly and conveniently measuring analytes by skilled laboratorian or by individuals who are unskilled as laboratorians. According to one or more embodiments of the invention, the fluorometer can include an optical block, a removable storage medium, an internal processor, a communications interface, and internal data storage.

15 The system and method generally comprises the fluorometer and a testing or assay device. The assay device is used in conjunction with the fluorometer to achieve a result regarding the concentration or presence of an analyte in a sample. Examples of analytes include chemicals, proteins, peptides, bacteria, viruses, nucleic acids, cellular organelles, cells, receptors and the like. The assay device can include reagents that are necessary for
20 performing an immunological or chemical reaction, such reaction giving rise to a change in fluorescence of the sample that has been treated with the reagents. The reagents can include chemicals, antibodies, peptides, analytes, analyte analogues and these reagents may or may not be coupled to fluorescent labels or to solid phases.

In one embodiment, the removable storage medium is implemented utilizing a
25 ROM chip or other memory device, which can be interfaced to the fluorometer to provide operating instructions as well as calibration curves and control and calibration data. Preferably, the memory device is mounted on a carrier which provides easy insertion and removal such that a plurality of memory devices containing specific sets of data can be easily interchanged. In this manner, the fluorometer can be easily programmed and re-
30 programmed to perform a variety of tests and calibrations.

Additionally, the removable storage medium can be implemented using a removable medium such as a disk and disk drive. The disk can contain test data sets for one or more types of tests to be performed. The test data sets can include test instructions

and calibration curves for the test, as well as other program information and calibration and control information for the instrument.

A communications interface can be included to facilitate communications between the fluorometer and one or more other devices. The communications interface can include
5 a wired and wireless interface to provide direct or networked communications. The communications interface can be used to download test data sets, including, for example, test identifications, test instructions and calibration curves, as well as other program information and calibration and control information. The communications interface can also be used to allow the fluorometer to share processing responsibilities with other
10 devices such as a computer or other processor. Such an interface (wired or wireless) can be implemented, for example, utilizing an RS-232, infrared or modem interface for direct connection, or a network interface for network communications to one or more processors.

In one operational scenario, the communications interface is used to allow a physician or other health care professional at a health care facility (e.g., a doctor's office,
15 clinic, testing center, hospital, or other health care site or facility) to transmit test instructions to the fluorometer with regard to which tests are to be performed for a particular patient. The interface can also be used to forward test results to a health care facility to apprise the health care professional of the results. Results of tests and a catalog of tests performed can be sent to various locations for patient-diagnosis, record-keeping,
20 billing, and other purposes.

In an alternative operational scenario, a patient can perform testing at home, and test results and instructions can be exchanged with a health care facility via the communications interface. In this embodiment, patients who require frequent monitoring can get the necessary tests without traveling to a health care facility each time a test is
25 needed, for example, as may be required by patients taking daily regimens of therapeutic drugs.

In yet another operational scenario, a technician in the field can measure water or ground contamination and transmit the results to a home office via a cellular telephone or other communications medium to inform officials of the progress of a decontamination
30 procedure.

Internal data storage can be used to store program instructions (including test instructions), calibration curves, control and calibration data as well as other data used in the operation of the fluorometer. Internal data storage can also provide register space for

operand storage. Internal data storage can be implemented using, for example, RAM or DRAM technology, or other memory technology. Disk or other storage space can be used to supplement the internal data storage, depending on storage cost and access latency tradeoffs. Cache techniques can also be used to optimize performance.

- 5 Data storage, either internal or removable, can be used to store test information regarding a test or tests conducted or to be conducted on one or more samples. The test information can include information such as, an identification of the patient and other patient information, a sample identification, an identification of a test or tests performed on the sample, a date and time at which the tests were conducted, test conditions, test
10 results, specific reagent information, such as lot numbers and expiration dates and other pertinent information. The test information can be stored in a record that can be indexed using, for example, the patient identification or other indexing designation.

- Various user interfaces can be provided to facilitate user control and to enhance operability of the fluorometer. Input interfaces can include data entry devices such as a
15 keyboard, keypad, touch-screen display, mouse, voice recognition input, or other data entry device. Output interfaces can include a display screen or monitor, printer, speaker or other output device.

- The assay mechanism according to one embodiment includes a motorized mechanism for transporting the assay device in the fluorometer. Examples of such a
20 motorized mechanism include, for example, those as described in U.S. Patents 5,458,852 and copending United States Patent Application No. 08/458,276, titled "Devices For Ligand Receptor Methods," which are incorporated herein by reference. The movement of the assay device in the fluorometer functions to position the diagnostic lane of the device with an optical block so that one or more fluorescent areas or zones of the assay device
25 can be measured. The degree or presence of fluorescence in the diagnostic lane is related to the concentration or presence of analyte in the sample. The optical block can include a light source, detector and optics used to excite the sample as well as to sense the fluorescence of the excited sample. In one embodiment, the sample is disposed on the assay device. The assay mechanism can provide the capability to transport the assay
30 device along the optical block such that fluorescence of one or more of a plurality of zones on the diagnostic lane of the device can be measured. As such, one advantage of the invention according to this embodiment is that enhanced testing algorithms can be utilized, if desired, in measuring the fluorescence of the sample.

An additional advantage of the invention is that the communications interface can be used to allow the fluorometer to be interfaced to networks such as, for example, a hospital or other health care facility network, or other information networks whereby the fluorometer can retrieve data which may be needed to conduct tests and download other data including the test results. Additionally, the communications interface can be used to interface with the fluorometer to a stand alone computer such as, for example, a personal computer or an office or a home office computer. In these configurations, the fluorometer can utilize the processing and peripheral capabilities of the stand alone computer or network resources to supplement its own processing and interface capabilities. In yet another configuration, the fluorometer can interface with an existing instrument that is interfaced to a network, such as, for example, an instrument in a hospital emergency department or critical care unit that dispenses medications for use by the hospital personnel. Interfacing the fluorometer to an existing instrument has advantages in that the interface of the fluorometer to the instrument can be one specific code, whereas the instrument interface code can be varied depending on the location of the instrument, for example, in different hospitals with different software interface codes.

For example, in one embodiment, the fluorometer can be operated as a portable hand-carried piece of test equipment that is used to test samples of blood. The portable, hand-carried unit can then be interfaced to a computer or computer network to upload test results or to simply communicate other data associated with the test and to use the processing power of the computer or computer network to perform some or all of the actual test processing. In yet another example, test data sets or other pertinent information can be downloaded from the outside entity to provide the fluorometer with guidance as to tests to be conducted on a particular sample. This guidance can be in the form of complete test instructions or simply an identification of a test to be performed for which the instructions are stored internally to the fluorometer. In another example, the data communicated to a network can be utilized in real time to diagnose and treat acute care patients.

Yet another feature of the invention is that it provides an encoded tag on the assay device such as, for example, a bar code label or magnetic strip to allow sample, test or reagent information to be encoded. Sample information can include, for example, an identification of the sample and sample type, an identification of the patient from which the sample was drawn, an indication of the test or tests to be performed for the sample, as

well as other data, as desired. Reagent information can include the type of reagents in a device, lot specific information, such as calibration information and expiration dating. Once a sample is correctly labeled, there is no longer a need for manual user intervention to enter this information. In fully automated embodiments, this information is stored along with test results and other pertinent information to create and maintain an accurate record of the tests and test results. As such, the chance for operator error in incorrectly identifying a sample or otherwise incorrectly entering information regarding a test, is minimized. Additionally, test results and other data relating to the test can be automatically stored along with the patient identification and other associated information such that data for a patient can easily be accessed.

Another embodiment of the invention is to utilize an encoder, such as for example a magnetic strip encoder, in the instrument to encode information on an assay device. For example, patient information, including patient number, tests to be performed and the like can be entered through the keypad of the instrument or via a centralized computer that downloads the information to the fluorometer. The encoder records the information onto the assay device, such that when the user inserts the assay device into the fluorometer, a reader reads the information on the assay device and combines the assay results with the encoded information. The combined information can be stored in the fluorometer and it can be communicated to a network for real time or later analysis.

Yet another feature of the invention is that internal data storage can be provided so that patient information and test results can be tracked in the form of a history log. For example, in a portable hand-held environment, a user or technician may test several samples of blood in a given time interval. The test results, along with the identification of the patient, can be stored in the local database such that a history log of tests and test results are maintained. This history log can then be downloaded via the communications interface or onto a removable media.

Still another feature of the invention is that in remote or in-home applications, the identification of the patient can be based on an automatic number identification (referred to as ANI). In this embodiment, when the patient's fluorometer dials the remote health care facility via a telephone network, the ANI signal provided by the telephone network is used by the health-care facility to identify the patient from which the communication originated. The ANI can be used in place of the an identification of the patient based on

the encoded label, or in addition thereto to provide a cross check against potential identification errors.

Further features and advantages of the invention as well as the structure and operation of various embodiments thereof are described in detail below with reference to
5 the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a diagram illustrating a representative functional architecture of a fluorometer according to one embodiment of the invention.

10 FIG. 2 is a diagram generally illustrating an operational scenario of the fluorometer according to one embodiment of the invention.

FIG. 3 is a diagram illustrating a representative functional architecture of the assay mechanism according to one embodiment of the invention.

15 FIG. 4 is a diagram illustrating an example implementation of the fluorometer with a health care facility according to one embodiment of the invention.

FIG. 5 is a diagram illustrating an operational scenario of the fluorometer in conjunction with a health care facility according to one embodiment of the invention.

FIG. 6 is a diagram illustrating an example implementation of the fluorometer in conjunction with a separate computer according to one embodiment of the invention.

20 FIG. 7 is a diagram illustrating a process for testing a plurality of sample areas according to one embodiment of the invention.

FIG. 8 is a diagram illustrating a process for removing a background signal from the test results according to one embodiment of the invention.

25 FIG. 9 is a diagram illustrating an example of spot locations in terms of signal strength according to one embodiment of the invention.

FIG. 10 is a diagram illustrating a process for integrating fluorescent intensity over the test range according to one embodiment of the invention.

FIG. 11 is a diagram illustrating an optical package used to excite the sample and receive the signal according to one embodiment of the invention.

30 FIG. 12, which comprises FIGS. 12A and 12B, is a diagram illustrating an example implementation of a chip carrier according to one embodiment of the invention.

FIG. 13 is a diagram illustrating in more detail one embodiment of a leading edge of the chip carrier illustrated in FIG. 12.

FIG. 14 is a diagram illustrating an example implementation of a ROM chip mounted on a printed circuit board according to one embodiment of the invention.

FIG. 15 is a diagram illustrating a cross-sectional view of a slot and tab according to one embodiment of the invention.

5 FIG. 16 is a block diagram illustrating an example computer system in which elements and functionality of the invention are implemented according to one embodiment of the invention.

DETAILED DESCRIPTION OF THE INVENTION

10 The present invention is directed to a system and method for performing fluorescent measurements including enhanced functionality. FIG. 1 is a diagram illustrating a functional block diagram of an enhanced fluorometer according to one embodiment of the invention. FIG. 1 illustrates examples of the functionalities that can be included with the automated fluorometer in terms of one example physical architecture, a
15 central bus structure. After reading this description, it will become apparent to one of ordinary skill in the art how to implement any or all of the described functionality using one or more alternative architectures.

 The enhanced fluorometer according to the embodiment illustrated in FIG. 1 includes a processor 104, a power supply 108, a user interface 112, a memory 116, a
20 communications interface 120, an assay device 124, a storage device 128, and removable storage media. In the example illustrated in FIG. 1, the removable media include a ROM chip 136 and ROM chip socket 132, and a disk drive 138. Any or all of these functionalities can be included with an enhanced fluorometer depending on the particular application.

25 Processor 104 controls the operation of the fluorometer and also provides control for the various functionalities provided with the fluorometer. Processor 104 can be a central processor that controls the functionality via a bus structure or other communications interface. Alternatively, for example, processor 104 can be implemented by distributing the processing functions among one or more of the various components
30 utilized to implement the functionalities of the fluorometer.

 Power supply 108 is used to provide necessary power to the fluorometer and its components. Power supply 108 can be implemented using batteries, solar cells, transformers used to convert an AC power source, or other techniques to provide the

appropriate power levels to the components. In one embodiment, power supply 108 is implemented using rechargeable batteries such as, for example, NiCad or Nickel Metal Hydride batteries which can be recharged using a charger connected to AC power from a conventional wall outlet. The use of rechargeable batteries provides a practical power source for portable applications.

User interface 112 provides one or more devices through which a user can interface with a fluorometer. In one embodiment, user interface 112 includes a keypad 162, a display 164, and a printer 166. Additional or alternative interfaces can be provided such as, for example, a keyboard, mouse, track ball, touch-screen display, or other user interface devices.

In one embodiment, keypad 162 is a small alphanumeric keypad that provides the user with input keys to assist in the direction of the functionality of the fluorometer. Keypad 162 can also include special function keys to perform single-touch operations. The function keys can be preprogrammed to perform specified functions, or user programs, depending on the application. Display 164 can be implemented using a number of display devices such as, for example, a small monochromatic LCD display. A small LCD display is preferable for portable applications because of its legibility and low power requirements. For non-portable applications display 164 can be implemented using, for example, a CRT or a color LCD display.

Likewise, printer 166 can be implemented using a variety of different printing techniques. For example, in the portable embodiment, printer 166 can be implemented using a small thermal printer such as that found on small calculators or adding machines. For non-portable applications, or applications where a portable fluorometer can be disconnected from the printer for mobility, larger printers can be used.

Memory 116 is used to provide storage for program data or other data used by processor 104 during operation. Memory 116 can be implemented using various RAM or ROM memory devices. Memory 116 can be used for example, to store operating instructions and to provide memory registers for operating and storage. Memory can also be used in conjunction with a storage device 128 such as a disk storage device. Storage device 128 can also be used to store program instructions, control and calibration curves, operational data, history logs, and other data which may be desired to be stored within the fluorometer. Preferably, storage device 128 is used to store large amounts of data, and the generally more costly but faster memory 116 is used to store only data which must be

accessed more frequently or more rapidly. A cache can be provided to minimize latencies associated with retrieving frequently used data from storage device 128.

ROM chip socket 132 can be included to provide a means by which a ROM chip 136 containing program instructions, calibration curves, control data, or other information
5 can be interfaced to the fluorometer.

Communications interface 120 can be provided to allow the fluorometer to communicate with various external devices. Depending on the desired applications and the environments in which the fluorometer is operating, various alternative communications interfaces can be provided. Communications interface 120 can be
10 implemented to include wired and wireless interfaces such as, for example, an RS-232 interface, an infrared interface, an RF interface, a network interface, or other communications interface appropriate for the application. Through the use of communications interface 120, the fluorometer can share information with other entities such as test results, test statistics, and other information, as well as receive information and
15 instructions from outside entities.

Assay mechanism 124 is used to perform the fluorometric readings on the sample in order to test the presence or concentration of one or more analytes. In one embodiment, assay mechanism 124 is a slide mechanism that is used to accept a small tray-like device, for example, an assay device. Assay mechanism 124 includes the optical components
20 necessary to perform the fluorometric readings as well as a slide on which the assay device slides to position the assay zones in the appropriate location so that fluorescence can be measured in a reproducible manner. In one embodiment, the mechanism is motorized such that the assay device can be automatically loaded and ejected from the fluorometer as well as positioned with respect to the optics during testing. In this embodiment where the
25 assay device is transported along a path in the slide, the path which includes optics used to excite the sample and sense the fluorescence. The path in which fluorescence is measured in the assay device is referred to as the diagnostic lane of the device.

In one embodiment, all of the data and instructions necessary to operate the fluorometer can be provided on ROM chip 136. In this embodiment, there is little or no
30 need for functionalities provided by memory 116 and storage device 128. In other words, in one embodiment, all of the memory requirements of the fluorometer are provided by the ROM chip 136. In alternative embodiments memory requirements are shared among or redistributed to any or all of these storage devices.

In one embodiment, a removable storage medium, such as, for example, ROM chip 136 or a disk in disk drive 138 is utilized to provide operating instructions to the fluorometer. In a preferred embodiment, this memory device is a ROM chip 136. The functionality of the fluorometer is described in terms of this preferred embodiment. After
5 reading this description, it will become apparent to one of ordinary skill in the art how to implement removable storage medium using other storage devices. In addition to operating instructions, ROM chip 136 can also be used to provide other pertinent data to the fluorometer to be used in controlling and calibrating the fluorometer as well as calibration curves for performing the various tests. In a preferred embodiment, ROM chip
10 136 includes test software used to conduct one or more tests. For example, test software can include program instructions used to direct the fluorometer to perform one or more fluorometric tests on a sample. Depending on memory space available on the ROM chip and test software size, one or more tests can be provided using a single ROM chip 136.

In one embodiment, each different type of test or assay is provided on a single
15 ROM chip. In this embodiment, each time a different type of test is desired to be performed, ROM chip 136 is replaced with the appropriate ROM chip 136 containing the desired test software in socket 132. In a preferred embodiment, the test software and associated calibration and control information, and software for a plurality of tests can be provided on a single ROM chip 136. In this preferred embodiment, the need to exchange
20 ROM chips 136 can be minimized. In a particularly preferred embodiment, the ROM chip 136 software is downloaded to the fluorometer so that a number of tests can be accessed by the fluorometer without changing ROM chip 136.

Where a plurality of tests can be provided on a single chip, user selection of those tests can be performed by user interface 112, or remotely via communications interface
25 120 or by bar code information on the assay device. An important functionality which can also be provided by ROM chip 136 is that different tests can be provided on different ROM chips such that "reprogramming" of the fluorometer to perform a variety of different tests can be accomplished simply by replacing ROM chip 136.

ROM chip 136 can also include calibration curves utilized to perform the desired
30 test. Because different tests typically utilize different calibration curves, in one embodiment, the calibration curves are provided on ROM chip 136 along with the test software. ROM chip 136 can also include control and calibration data to calibrate the fluorometer using controlled solutions. Because control and calibration information may

change based on the test being performed, this information in one embodiment is provided with each ROM chip 136 such that the fluorometer can be properly configured and calibrated to perform the desired tests.

ROM chips 136 containing instrument specific, test specific, and calibration specific information can be the same or different. In a preferred embodiment, specific chips are used to provide specific functionality. Examples of this functionality are now described according to one embodiment of the invention. Instrument specific operational software for one or more tests resides on a single ROM chip 136 termed a program ROM key. A chip termed a reagent code chip provides test specific information, including calibration information for one or more tests and lots of tests. Control solution information, including concentrations and ranges of analytes and expiration dating of the solutions resides on another ROM chip 136 termed QC Sample Code Chip. Calibrator solution information, including concentrations and ranges of analytes and expiration dating of the calibrator solution resides on another ROM chip 136 termed Calibrator Code Chip. Information relating to the accessibility of a user to change fluorometer parameters, such as addition and deletion of user passwords, normal ranges for the analytes being measured, frequencies of measuring control solutions and the QC simulator, and the like on yet another ROM chip 136 termed Supervisor Code Chip. Additionally, information relating to expected values measured by a QC simulator resides on yet another ROM key 136 termed Instrument Validation Code Chip.

FIG. 2 is an operational flow diagram generally illustrating an operation of the fluorometer according to one embodiment of the invention. In a step 204 software is loaded to perform the desired test. In one embodiment, as described above, a ROM chip or memory device 136 is inserted into socket 132, the ROM chip 136 containing the software for the test to be performed. Alternatively, the test software can be loaded using communications interface 120, disk drive 138, ROM chip 136, storage device 128 or other interface. The loading can include the transfer of any or all instructions and data to internal memory 116 or storage 128, or can simply encompass making these instructions and data available to processor 104 via access of the utilized device or interface.

For a new type of test, or at periodic intervals during testing, it may be desirable to calibrate the fluorometer. As such, in a step 208, the fluorometer is calibrated. As disclosed above, the calibration can be performed using calibration software with controlled test solutions. In one embodiment, an assay device, termed a QC simulator,

comprising a fluorescent zone or zones is inserted into the fluorometer. The fluorescent value of the QC simulator is a known value that can be loaded into the fluorometer via a ROM key or a bar code on the assay device. In another embodiment, a fluorescent chip, termed an internal calibrator, is within the fluorometer that is read by the fluorometer prior to each measurement of an assay device. The intensity of the fluorescence of the internal calibrator is a known quantity.

In a step 212, an assay device with sample to be tested is loaded into the fluorometer. In one embodiment, this is accomplished by inserting an assay device containing the sample into assay mechanism 124. In one embodiment, a bar code symbol or other encoded tag is used to provide an identification of the sample to be tested. In this embodiment, the encoded tag is read by the fluorometer such that the test results can be correlated with or later associated with the particular sample being tested. Examples of an encoded tag can include a bar code symbol, an encoded magnetic strip, a character designation capable of being read by an optical character reader, or a tag made using some other encoding technique.

In a step 216, the sample is tested. Samples to be tested can include, for example, biological fluid such as blood, serum, plasma, urine; a fecal extract and the like; an environmental sample such as water, a ground extract, a chemical and the like; or an extract of a food product. The testing is performed in accordance with the instructions provided by the software loaded in step 204. When an assay device is used as the carrier for the sample, various testing techniques can be used to optimize the test results. Examples of such techniques relating to immunological reactions are described in Principles and Practice of Immunoassay, C.P. Price and D.J. Newman, Macmillan Reference Ltd., 1997, and U.S. Patents 5,028,535, 5,089,391, 5,143,852, 5,458,852, 5,480,792 and 5,525,524 which are incorporated by reference herein. A preferred embodiment is used in conjunction with a motorized assay mechanism 124 to allow a plurality of sample zones or regions on an assay device to be tested.

In a step 220, the results of the tests are provided. The test results can be printed on printer 166, displayed on display 164, stored in a local memory or storage device within the fluorometer, written to a medium such as a disk in disk drive 138, or communicated to an external entity via communications interface 120. Preferably, the test results are displayed, printed, stored or transmitted along with the identification of the sample obtained using the encoded tag such that the test results can always be associated

with the proper sample. The use of an encoded tag in this manner helps to assure that the test results are always associated with the proper sample.

FIG. 3 is a diagram illustrating an example implementation of an assay mechanism or assay device drive according to one embodiment of the invention. The assay device drive according to the embodiment illustrated includes drive electronics 304, a position encoder 308, and an encoded tag reader 312 such as, for example, a bar code reader. In one embodiment, drive electronics 304 includes a motor to position the assay device and a motor controller to control the motor. A friction drive, gear drive, or other mechanism can be used to translate the rotation of the motor into motion of the assay device. Drive electronics 304 are thus used to load and eject the assay device as well as to position the assay device with respect to the optics of the fluorometer, for example, along the diagnostic lane. In this embodiment, the assay device is moved in relation to stationary optics. In alternative embodiments, the optics can be moved instead of, or in addition to, the assay device.

Position encoder 308 is used to determine the position of the assay device within assay device drive 300. Position encoder 308 can obtain position information from the assay device itself such as, for example, by sensing an encoded label on the assay device. Alternatively, position encoder 308 can determine the position of the assay device based on the rotation of the drive shaft through the motor using well-known encoder techniques. Encoded device reader 312 is used to read the encoded tag provided on the assay device. In one embodiment, encoded tag reader 312 is a bar code reader that reads a bar code label on the assay device. Alternative embodiments can include, for example, a magnetic stripe reader, an inductive reader, or an optical character recognizer. An encoded tag reader 312 senses the encoded tag information from the label on the assay device and provides this information to processor 104. The encoded information can include information such as, for example, a patient I.D., an identification of the tests to be performed on the sample, an identification of the sample type, or other appropriate or pertinent information. This information can be used to log the test results as well as to control the type of testing performed or test parameters used.

In one embodiment, drive electronics 304 and position encoder 308 are used to control the positioning of the assay device, as well as to reposition the assay device during testing such that a plurality of regions of the assay device can be tested. This capability to position the assay device such that various portions of the sample can be tested allows

enhanced testing algorithms to be utilized to produce improved test results. An example of enhanced testing routines that can be used where different regions of an assay device are tested is fully described in copending patent applications of common assignee, having Application No. 08/311,098, and titled "Fluorescence Energy Transfer and Intramolecular Energy Transfer in Particles Using Novel Compounds" and Application No. 08/409,298 also titled "Fluorescence Energy Transfer and Intramolecular Energy Transfer in Particles Using Novel Compounds," which are incorporated herein by reference.

FIG. 4 is a block diagram illustrating an example implementation of a fluorometer 100 used in communication with an information system 408 to perform fluorometric tests.

As described above, a communications interface 120 can be provided to allow the fluorometer 100 to communicate with outside entities such as, for example, a hospital network, a physician's office, a testing clinic, other laboratory computers, or other relevant entities. To illustrate the utility of the communications interface, a simple example scenario is now described. In this example scenario, the outside entity is a health care information system 408 such as, for example, a hospital data system that can be accessed by a physician or other health care professional to order tests for and treat a patient.

In the example illustrated in FIG. 4, information system 408 is implemented to include a data entry terminal 462 and a data storage device 464. After reading this description, it will be apparent to one skilled in the art how information system 408 can be implemented using alternative architectures. In one example scenario, a blood sample is drawn from the patient. The vials containing the blood are labeled with an identification of the patient and sent to the lab that will perform the fluorometric tests. The technician at the lab receives the blood sample, prepares the assay device and places the encoded I.D. label 412 onto the assay device. I.D. label 412 identifies the assay device as belonging to the patient from which the sample was withdrawn. I.D. label 412 can be generated by the lab technician or transferred from the received vials.

FIG. 5 is an operational flow diagram illustrating an example process by which the sample can be tested in an automated fashion with fluorometer 100 in communication with information system 408 utilizing communications interface 120. In a step 502, an assay device is prepared with a sample and labeled with an encoded tag. In a step 504, the prepared sample, that is, an assay device with the sample added, is loaded into fluorometer 100. In one embodiment, the assay device is a cassette or tray-like container. The assay device includes an encoded I.D. tag 412 which can provide, for example, an identification

of the patient from which the sample is drawn, an indication of the assay or assays to be performed and can include additional information as may be deemed relevant for the particular application. Any additional information required can be downloaded or entered by the health care professional via user interface 112.

5 In a preferred embodiment, the encoded I.D. tag 412 provides an identification of the type of test that is being performed, for example, a test that assesses the condition of a patient with respect to a myocardial infarction. In this preferred embodiment, the lab technician enters the patient information, for example, the patient number, via keypad 162. In yet another embodiment, an encoded label is not used and all necessary information is
10 obtained through downloading, manual entry or a combination thereof.

 In a step 508, fluorometer 100 reads the encoded tag on the assay device. In a step 512, information from encoded tag 412, and/or any additional information downloaded and/or entered, or a translation thereof, is sent to information system 408 to access information or to download information about the patient or about tests ordered for that
15 patient.

 For example, the health care professional at the hospital ordering tests for the patient can enter in the patient's name or other identification as well as a list or identification of tests ordered for that patient into system 408. The patient's name or identification and the tests to be performed for that patient are stored in database 464 so
20 that there is a record of the tests ordered for that patient. Thus, in step 512, the information sent to information system 408 is used to access the database, to thereby retrieve information indicating which tests are ordered for the patient. In a step 516 test instructions which can include, for example, instructions for performing a desired test, or an identification of the test or tests ordered for the patient, are downloaded to fluorometer
25 100 using communications interface 120.

 In a step 520, fluorometer 100 performs the test, after the technician loads the assay device, by either carrying out the downloaded instructions, or by accessing the instructions locally (e.g., by local storage, by input from keypad 162 or from ROM chip) based on an identification of the test received from information system 408. Where a
30 different ROM chip 136 or disk needs to be installed to perform the required test, the user is informed via user interface 112.

 In a preferred embodiment, the testing is performed automatically without user intervention. Indeed, in one embodiment, timing algorithms such as those described

below are implemented to control the timing of the assay process. In this embodiment, the test is carried out and completed automatically without intervention. Alternative embodiments may provide the capability of the system to prompt the user for certain inputs as deemed appropriate.

- 5 Once the tests are completed, in a step 524, the results are provided to information system 408. The results can be stored in database 464 and are identified as being associated with the patient. As such, results of the tests can be accessed by health care professionals using terminal 462 and can be printed to provide a hard copy of the results.

- Additional information regarding tests conducted on the sample can be transmitted
10 to information system 408 as well. The additional test information can include, for example, an identification of the patient and other patient information, a sample identification, an identification of a test or tests performed on the sample, a date and time at which the tests were conducted, test conditions, test results and other pertinent information. This test information can be used to update database 464 such that
15 information system 408 has a complete record of tests, results and associated data for a patient. Additionally, the test information can be stored in the fluorometer in local storage 128, memory 116 or on removable storage medium (i.e., ROM chip 136, removable disk, etc.).

- In order to assure that all required tests are performed, and also to avoid
20 duplication of testing, record flags or other techniques can be used when the database 464 is accessed to retrieve test instructions. For example, when fluorometer 100 accesses information system 408 to receive instructions for a particular test, that test is flagged as being performed such that subsequent accesses by this or another fluorometer 100 will not retrieve the same test instructions. Once a test is completed and the results provided to
25 information system 408, another flag can be set indicating the status of the test as being completed.

- Note that where tests are ordered which depend on the type of sample being tested, I.D. 412 can also include a sample identification or a sample type identification. For example, where a prepared sample is plasma, an indication of such is included on I.D. 412,
30 or alternatively entered by the user using the keypad or other input device on fluorometer 100. Thus, when fluorometer 100 queries information system 408 to access test instructions, tests which are ordered for plasma are checked, and if present, retrieved from database 464. As illustrated by this example scenario, the chance for operator error in

identification of the sample and performing the appropriate tests on that sample, are minimized at the fluorometer. In this scenario, once the sample is loaded with the correct I.D., tests ordered by the physician or health care professional at the hospital are electronically retrieved, automatically performed, and electronically reported back to information system 408. As such, there is little or no need for human intervention in this system, with the exception of addition of the sample to the assay device and insertion into the fluorometer by the technician.

In yet another example embodiment, label 412 can be used to further automate the testing process, even in a stand-alone environment. For example, in one embodiment, when the technician prepares the blood sample, the label 412 code includes a description or indication of the test to be performed on that sample. When the fluorometer reads the test description or indication, the appropriate test is performed. As with the remote example described above, the test can be automatically accessed from local storage or the user may be prompted to load the test-related information via ROM chip 136, disk or other memory or storage device. Information from the tag 412 as well as tests performed and their results can be used to create a record locally on fixed or removable media as well as remotely via communications interface 120. Information from the record can be used for patient diagnosis, accounting, billing, statistical and other purposes.

As would be apparent to one of ordinary skill in the art after reading the above descriptions, there are numerous scenarios that can be implemented to take advantage of the capabilities of fluorometer 100 utilizing an encoded I.D. label 412 and communications interface 120. Also, as described above, communications interface can take on a variety of different physical embodiments such that communication between fluorometer 100 and information system 408 can be implemented using a wireless communication link, a hardware communication link, networks communication, or other communication facilities.

As an example alternative application to that described above with reference to FIGs. 4 and 5, consider an application in which a patient utilizes fluorometer 100 to conduct in home testing. In this application, the patient obtains a sample and places it in the assay device or cassette. The sample can be a blood sample, for example, from a finger prick, a urine sample or other appropriate sample.

Fluorometer 100 can be pre-loaded with test instructions appropriate to the particular patient. Alternatively, fluorometer 100 can utilize communication interface 120

to access test instructions from the clinic, physician's office, hospital or other health-care facility. The patient loads the sample into fluorometer 100 and the appropriate test or tests are conducted.

Test results are forwarded to the health care facility using communications interface 120 so that the patient's records can be updated and the appropriate health care professional can be apprised of the results.

Fluorometer 100 can include an alarm feature that automatically informs or reminds the patient when a test is required. This feature can be implemented, for example, using a built-in programmable clock. The clock can be programmed manually or by installing scheduling instructions. For example, a scheduling program detailing the type or types of tests to be performed and the time for such tests can be installed before fluorometer 100 is delivered to the patient. The installation can be performed, for example, using removable storage media. Schedule programming can also be downloaded via communications interface 120 both initially, and as treatment progresses. For example, the tending physician may wish to update the testing schedule by adding new tests, changing the testing interval, or discontinuing certain tests. The physician can update information system 408 to reflect this change and the changes can be downloaded via communications interface 120. In this manner, the testing can be tailored and updated to suit a patient's current needs.

In one embodiment, the scheduling program indicates the type of test to be performed at a scheduled test occurrence. Information regarding the test type can be, but does not need to be provided to the patient. This information, however, along with test results, is preferably provided via communications interface 120 to update the patient's data base or to inform the physician of the test results. The scheduling program can also indicate the sample type required for a particular test.

In embodiments where the communications interface is a telephone, the patient's identification can be provided to the health-care facility using automatic number identification, or ANI, based on the telephone number from which the test instrument is calling. Where ANI is not available, or where the patient is calling from a location where his or her ANI is not recognized, the system may prompt the patient to enter an identification.

In yet another embodiment, communications interface 120 can be utilized to inform the patient that a personal visit to the health-care professional is required. For

example, consider a scenario in which the tending physician is obtaining and reviewing results of regularly conducted tests. The physician may detect a change in condition that warrants a personal appointment. The physician can input this information into the health care information system and the patient is apprised of the necessary appointment.

5 In one embodiment, the health care information system maintains a calendar of available appointment slots for the tending physician. The appointments can be maintained on the physician's personal computer or on a data server or other accessible data base within the health care information system to facilitate sharing of calendar information among health care professionals.

10 A listing of available appointments is sent to the patient via communications interface 120. The patient, using user interface 112 reviews the available appointment slots and selects one that fits his or her schedule. Fluorometer 100 provides this information to health care information system to reserve that slot for the patient. A reminder for the appointment can be programmed into the internal clock of fluorometer
15 100 to remind the patient a desired time interval before the appointment. Alternatively, the health care information system can store this information and send an appropriate reminder to the patient via communications interface 120.

The use of communications interface 120 in the in-home embodiments allows real-time or near-real-time interaction between the patient and the tending health-care
20 professional although the patient and health care professional are not at the same location. Tests can be conducted and the results reported and analyzed as they are performed. Test schedules can be updated and otherwise modified as the health-care professional deems fit based on results of recent tests.

As described above, the fluorometer can be utilized in conjunction with a separate
25 processor such as, for example, a workstation or personal computer. In this embodiment, a variety of fluorometer functionalities can be delegated to or shared with the separate processor. To further illustrate this capability, it is now described in terms of an example scenario. In this example scenario, fluorometer 100 is interfaced to a personal computer 604 as illustrated in FIG. 6. In one embodiment, the interface is implemented using an
30 RS-232 communications interface, although other communications interfaces can be selected based on speed, cost and reliability tradeoffs.

As an example of the manner in which a processor such as a personal computer 604 can be utilized to enhance the performance of fluorometer 100, consider a scenario

where fluorometer 100 performs a plurality of tests on blood samples. After performing the appropriate measurements and readings, the fluorometer provides these results to personal computer 604. In a sense, fluorometer 100 can be considered as providing raw data to personal computer 604 although a certain amount of processing can be performed by fluorometer 100. Personal computer 604 accepts the raw data and performs analysis and processing of the data to help to arrive at definitive test results or to interpret therapy for the patient based on the test results. Test results and related data can be described alphanumerically and graphically on the display screen of personal computer 604. Through user interfaces on personal computer 604 an operator can adjust the functionality of the test as well as test parameters to obtain optimum results.

As disclosed above, in one embodiment the assay device is repositioned during testing such that different areas of the sample can be excited and the fluorescence of each of these areas can be measured. An example process by which this is accomplished and the data analyzed now follows. FIG. 7 is an operational flow diagram illustrating this process according to one embodiment of the invention. In a step 704, the assay device is stepped along the diagnostic lane such that at each step a different region of the assay device is excited and fluorescent measurements are sensed from that region. In one embodiment, the assay device is stepped in only along one direction (e.g., back and forth or side-to side) and therefore the regions or zones are divided along one direction of the assay device. In alternative embodiments, the assay device can be stepped, for example, both back and forth, and side to side, such that additional regions on the assay device can be defined.

In a preferred embodiment, the assay device is stepped along the diagnostic lane in discrete steps, and fluorescence measured at each step. In alternative preferred embodiments, the assay device is driven continuously at a given speed along the diagnostic lane, and measurements are taken throughout the process. In either embodiment, the measurements themselves can be at discrete intervals or continuous.

In a step 708, the results of each region are recorded. In embodiments where the device has a fluorescent result, the recorded results represent optical amplitudes sensed from the excited region. Preferably, the data from a particular sample are recorded and processing is performed upon recorded data such that processing does not have to be performed in real time, and the entire data sample can be utilized in processing.

In a step 710, data processing begins with checking the quality of the trace. The details of this step according to one embodiment are fully described in copending patent applications of common assignee, having Application No. 08/311,098, and titled "Fluorescence Energy Transfer and Intramolecular Energy Transfer in Particles Using Novel Compounds" and Application No. 08/409,298 also titled "Fluorescence Energy Transfer and Intramolecular Energy Transfer in Particles Using Novel Compounds."

As the assay device is stepped along and moved in relation to the optics, nonspecific binding of the fluorescent label results in a residual fluorescent signal. This signal is a background signal that can be highly variable from sample to sample.

Therefore, in a step 712 it is desired to remove this signal before calculating the amount of fluorescence immobilized on each area of the diagnostic lane for each sample tested. The fluorescence intensities measured for each area or region tested are integrated to determine the total amount of fluorescent label immobilized as a result, for example, of an immunological binding reaction.

In a step 716, the processed signal without the background is integrated to determine the total fluorescence of the sample. In a step 718 the result is computed, for example by comparison to a threshold, to determine whether the test is positive or the concentration of each analyte being tested.

In fluorometer embodiments, the sensed amount of fluorescent label is a function of the acquisition method. More specifically, the sensed amount of fluorescent label for a given region is the product of the fluorescent strength emitted by that region and the width of the region. Therefore, if the displacement of the assay device varies from the expected displacement the results obtained will be inconsistent and will not be reproducible. For example, if the displacement is shorter than expected, the amount of fluorescent label in the region will be reported as higher for a given sample.

To compensate for changes or inconsistencies in positioning, position encoder 308 can be utilized to obtain a measurement of the actual or relative displacement during the test process. In a preferred embodiment the position is encoded by monitoring the rotation of the motor, i.e. using a rotary position encoder. After reading this description, it will become apparent to one of ordinary skill in the art who to implement this functionality utilizing alternative embodiments of rotary encoders. From this, errors in the expected displacement arising from changes in motor speed are factored out.

However, imperfections in the drive mechanics and variations in the load on the drive mechanics can result in errors between the measured motor position and the actual assay device position. In a preferred embodiment the average error between the measured motor position and the assay device position is recorded as a function of the position of the assay device. This information is used to map the measured motor position into actual assay device position. In a particularly preferred embodiment the position is encoded by monitoring the position of the assay device with respect to the optics, i.e. using a linear position encoder. After reading this description, it will become apparent to one of ordinary skill in the art who to implement this functionality utilizing alternative embodiments of linear encoders. As a result of this process, inconsistencies in motion can be factored out to obtain a more accurate and reproducible test result.

FIG. 8 is an operational flow diagram illustrating a manner in which the background signal can be removed according to one embodiment of the invention. In a step 804, the spot locations or zones of fluorescence as a result of the assay process are determined by where a signal is received. Alternatively, in a preferred embodiment, spot locations or assay zones are defined by the fluorometer software such that the fluorometer is programmed to measure fluorescence in a particular location or locations along the diagnostic lane of the assay device. These spot locations or zones are the points along the length of the diagnostic lane of the assay device at which an amplitude above a determined threshold is received.

FIG. 9 is a diagram illustrating an example of spot locations along the length of the diagnostic lane. In FIG. 9, the ordinate is the amplitude of the fluorescent signal received, and the abscissa is the length along the diagnostic lane. Because the assay device is moving along the diagnostic lane, this also represents the position of the assay device with respect to the optics. Thus, travel along the abscissa represents a position in the diagnostic lane along the assay device. As illustrated in FIG. 9, there are areas where it is apparent that a signal is received as illustrated by spot 920, separated by areas in which there is merely background noise 914. Areas of apparent signal readings are referred to as spot locations or (assay) zones 920.

In a step 808, the level of background signal 914 on either side of a spot 920 is determined. In a step 812, the level of background signal from one side of the spot to the other side is assumed to have a constant variation. This level is considered the background and is subtracted from the signal level of the spot location in a step 816. As a

result of this process, a representation of the actual signal level without the contribution of the background noise can be determined. FIG. 10 is a diagram illustrating a process by which the fluorescence intensity is integrated over each spot 920 according to one embodiment of the invention. In a step 1002, the sum of the adjusted fluorescent intensity is calculated.

In a step 1004, this intensity is multiplied by the step distance between each point. The step distance relates to the amount by which the assay device is moved along the diagnostic lane between each data reading. In one embodiment, variations in the step distance are factored out of the readings in step 1004 by correcting the step distance according to previously measured calibration data.

In a step 1006, this result is normalized by the appropriate meter gain. The meter gain (G) is a scale factor that represents the relative intensity of the excitation source and the relative sensitivity of the detector. The detected signal is the product of the source intensity, the amount of fluorescence and the sensitivity of the detector. In one embodiment, the result is normalized by the width of the spot and therefore relates to the average amount of fluorescent label as a result of the assay process. In a preferred embodiment, the result is not normalized by the width of the spot and therefore relates to the total amount of fluorescent label as a result of the assay process.

The source is subject to both frequency and intensity drifts. These variations are tracked jointly with a fluorescent spot in the meter with known response at the nominal power and wavelength of the source. Therefore, the fluorescent signal of the test zone will scale proportionally to the variation in measured signal from the internal standard. One skilled in the art will recognize that the internal standard will correct any fluctuations in the sensitivity of the detector as well. In a preferred embodiment, step 1006 also normalizes the result to the measured signal of the internal standard.

In a preferred embodiment, the intensity (I_j) at each region is measured by subtracting the baseline signal (BL) from the measured signal (MI), $I_j = MI_j - BL_j$, as illustrated in FIG. 8. The intensity of each region is then multiplied by the length of each region, e.g. the known distance between measurements. The length of each region, ΔX_j , can be a function of position along the diagnostic lane and is not necessarily a constant.

The sum of this product for each region within a zone represents the uncorrected fluorescent label (UFL) within the zone ($UFL = \sum I_j \Delta X_j$). The UFL is multiplied by the meter gain, which scales the result to be independent of the meter. The UFL is also

multiplied by the expected internal standard value (EISV) and is divided by the measured internal standard value (MISV), yielding an instrument independent result of the total fluorescent label (TF).

Conventional fluorometers used to measure fluorescence of a sample typically operate in the visible or near-visible wavelength range. These assay methodologies utilize dyes that fluoresce, generally between 400 nm and 500 nm. Such dyes include but are not limited to fluorescein and methyl-umbelliferyl phosphate. However, blood, plasma and serum samples themselves absorb and fluoresce in the ultraviolet and visible spectrum, up to about 600 nm. As a result, the signal-to-background ratio of the signal sensed from the fluorescent dye is eliminated or diminished.

To overcome these difficulties, a preferred embodiment of the invention utilizes dyes that excite and emit in the infrared or near-infrared range, particularly between about 600 nm to 1300 nm. As a result of these characteristics, certain advantages can be gained. First, the excitation wavelength of the dye is not a wavelength that, for example, the blood or serum absorbs. As such, energy from the excitation source is not lost. Second, the emission wavelength does not correspond to the absorption of the sample such that fluorescent light is not lost. Finally, because the blood and serum do not fluoresce at these wavelengths, there is reduced background noise sensed by the detector.

Another preferred embodiment makes use of dyes that possess Stokes shifts greater than about 90 nm. Stokes shifts of greater than 90 nm allow simplification of the design of the optical block, in that special band pass filters are not necessary to block excitation light energy. In other words, the overlap of excitation and emission light is minimized as the Stokes shift increases, thus increasing the recovery of fluorescent light from assay device. One skilled in the art recognizes that the overlap of wavelengths of excitation and emission light requires the use of band pass filters to prevent measuring the excitation light when it is desirable to measure the emitted light. The use of band pass filters has the disadvantages of cost and of decreasing the yield of measured fluorescent light.

Dyes that exhibit properties of excitation and emission wavelengths in the near infrared and infrared and have Stokes shifts greater than about 90 nm are fully disclosed in copending United States Patent Application Serial Number 08/601,492, titled "Fluorescence Energy Transfer in Particles," Application Serial No. 08/274,534, titled "Fluorescence Energy Transfer in Particles," Application Serial No. 08/311/098, titled "Fluorescence Energy Transfer and Intramolecular Energy Transfer in Particles Using

Novel Compounds,” and Application Serial No. 08/409,298, titled “Fluorescence Energy Transfer and Intramolecular Energy Transfer in Particles Using Novel Compounds,” which are incorporated herein by reference. A preferred embodiment utilizes dyes and dye systems of phthalocyanines and hybrid phthalocyanine derivatives that incorporate fluorescence energy transfer in particles, particularly in latex particles having bound antibodies, proteins, ligands and ligand analogues.

FIG. 11 is a block diagram illustrating an example implementation of an optical package that operates effectively with dyes exhibiting these desired properties according to one embodiment of the invention. The optical package illustrated in FIG. 11 includes an energy source 1102 and an energy detector 1104 (i.e., an optical transmitter and receiver). Additionally, cut-off or band-pass filters 1108, 1110 can be provided to filter background signal from the energy source 1102 or other sources.

Energy source 1102 is preferably an optical energy source 1102 that emits light in the infrared or near-infrared range portion of the spectrum, but can also emit light in the ultraviolet or visible wavelengths. In a preferred embodiment, optical energy source 1102 emits energy at a wavelength of approximately 670 nm. This wavelength of energy is poorly absorbed by blood or serum and does not cause blood or serum to fluoresce, at least not appreciably. Preferred energy sources are flash lamps, light emitting diodes and laser diodes. Particularly preferred energy sources are laser diodes.

Depending on the dye used, the fluorescent energy emitted from the sample is at a wavelength different from the excited energy. Specifically, in a preferred embodiment, the wavelength of the energy emitted as a result of the fluorescence of the dye is at approximately 760 nm. This wavelength is sufficiently different from that of the excitation wavelength (670 nm), such that energy (excitation) source 1102 will not significantly contribute to the background signal measured by the energy detector 1104. In a preferred embodiment, a cutoff filter is used to minimize light greater than about 690 nm from the source.

To further improve the signal-to-background ratio of the system, one or more filters can be included. In a preferred embodiment, for example, a high-pass filter 1108 is used to cutoff frequencies of the energy source at frequencies below (wavelengths above) the preferred excitation wavelength which would potentially be a source of background for energy detector 1104. Additionally or alternatively, a low-pass filter 1110 can be included to cutoff frequencies of the energy source at frequencies above (wavelengths below) the

preferred detection frequency. Preferred energy detectors are photomultiplier tubes and silicon photodiodes. Particularly preferred energy detectors are silicon photodiodes.

In one embodiment, energy source 1102 and energy detector 1104 are positioned such that the excitation energy emitted from energy source 1102 impinges on the sample at approximately a 45 degree angle. Alternative configurations can be implemented using different angles. Additionally, a plurality of energy sources 1102 and/or detectors 1104 can be implemented to optimize the energy readings. One skilled in the art will recognize that sources generally do not provide uniform illumination, and furthermore, a 45 degree angle of incidence will result in a variation in illumination across the sample. In a preferred embodiment, source 1102 contains a special optic that provides for homogenous illumination of the sample. In a preferred embodiment this special optic is a micro-lens array. In a particularly preferred embodiment the special optic is a diffractive optic, which provides homogenous illumination, including correction for the 45 degree angle of incidence.

Because the output of light sources such as laser diodes often have non-uniformities, inconsistent results may be obtained from one light source to the next. To account for these non-uniformities a flattener can be used in one embodiment to achieve a more uniformly distributed beam pattern. Examples of such flatteners can include diffusers and diffractive optics.

As described above, a removable storage medium can be included to facilitate or enhance the operation of a fluorometer. Specifically, in one embodiment described above a removable storage medium is implemented using a socket 132 and a ROM chip 136. Preferably, in this embodiment, ROM chip 136 is mounted in a chip carrier that includes contacts to interface with socket 132. In this embodiment, the chip carrier can be inserted into and withdrawn from socket 132 to facilitate the change or swapping out of ROM chips 136. An example of ROM chip carrier is now described.

FIG. 12, which comprises FIGS. 12A and 12B, is a diagram illustrating an example implementation of a chip carrier 1200 according to one embodiment of the invention. FIG. 12A is a perspective view of what is referred to as the top surface of chip carrier 1200. As illustrated in FIG. 12A, chip carrier 1200 includes a body portion 1203 and a rear tab 1209. Body portion 1203 includes a structure for mounting a ROM chip 136, as well as for guiding ROM chip 136 into socket 132. Rear tab 1209 provides a tab-

like structure that facilitates the handling of carrier 1200 and the insertion and removal of carrier 1200 into and out of socket 132.

In the embodiment illustrated in FIG. 12, carrier 1200 includes a top face 1204 that has an open area 1205. Open area 1205 provides an opening through which contacts in socket 132 can access electrical contacts 1408 (illustrated in FIG. 14). Lateral members 1206 define the sides of carrier 1200 and form boundaries of open area 1205. A central member 1206 can be included to provide additional structural rigidity, as well as to assist in the guidance of carrier 1200 in socket 132. Leading edge 1208 of carrier 1200 is preferably beveled, as are lateral members 1206. Because these members are beveled, they include a plurality of edges and faces that facilitate alignment during insertion as well as facilitate a tight fit of carrier 1200 within socket 132. Leading edge base 1212 provides structural rigidity for lateral members 1206 and central member 1207. FIG. 12B is a diagram illustrating a bottom view of carrier 1200 according to one preferred embodiment of the invention. As illustrated in FIG. 12B, lateral members 1206, cross member 1207 and leading edge base 1212 provide a frame around a chip cavity 1236. It is chip cavity 1236 in which ROM chip 136 is disposed. In one preferred embodiment, ROM chip 136 is mounted on a relatively flat structure such as a printed circuit board. An example implementation of this embodiment is illustrated in FIG. 14. As illustrated in FIG. 14, ROM chip 136 is mounted on a printed circuit board 1404. A printed circuit board 1404 includes a plurality of contacts 1408 that are used to make electrical contact with corresponding contacts in socket 132. Leads 1406 are used to connect the leads of ROM chip 136 with contacts 1408.

In one embodiment, contacts in socket 132 (not illustrated) are implemented using wiper contacts similar, for example, to those found in phone jacks or circuit board card-edge connectors. To maintain solid electrical contact, the wiper contacts can be spring loaded using, for example springs, or the sprung force of the bent contact metal itself. The contacts in socket 132 as well as contacts 1408 and leads 1406 are implemented using a conductive material, such as, for example, copper, gold, silver or other conductive material.

The width and thickness of contacts in socket 132, leads 1406 and contacts 1408 as disposed on printed circuit board 1404 can be varied to provide the proper amount of conductivity depending on the implementation. Additionally, the area and thickness

provided for a ground plane or ground lead 146 is chosen to provide adequate current handling capacity for grounding.

Printed circuit board 1404 in one embodiment includes tabs 1412. Tabs 1412 align with recesses 1214 to facilitate placement of ROM chip 136 and circuit board 1404 within chip cavity 1236. A flexible tab 1234 holds printed circuit board 1404 in place. A crenelated or other structure 1242 can also be provided to help hold circuit board 1404 in place. Structure 1242 can be crenelated as illustrated in FIG. 12B, a contiguous structure, or some other alternative structure. The combination of structure 1242 and tab 1234 allows circuit board 1404 to be slid under structure 1242 and snapped into place below an edge of tab 1234. A ridge 1244 extending around or partially around the interior of chip cavity 1236 provides a support on which printed circuit board 1404 can rest. The combination of ridge 1244 in conjunction with structure 1242 and tab 1234 holds circuit board 1404 firmly in place within chip cavity 1236. In one embodiment, a recess 1237 is provided as an artifact of the molding process.

Chip carrier 1200 can be implemented using a molded acrylonitrile butadiene styrene polymer (ABS), a polyoxymethylene (POM) polymer, a styrene polymer or styrene copolymers. In one embodiment, tab 1234 extends from slot 1210 to facilitate the flexibility of tab 1234 for insertion and removal of printed circuit board 1404. FIG. 15 is a diagram illustrating a cross-sectional view of slot 1210 and tab 1234. As illustrated, in this embodiment, tab 1234 is molded as portion of top base 1204. The presence of slot 1210 provides additional flexibility to tab 1234 without stressing the area where tab 1234 meets top surface 1204. Also illustrated in FIG. 15 is the interface between tab 1234, the bottom surface of top face 1204, and printed circuit board 1404.

FIG. 13 is a diagram illustrating in more detail one embodiment of leading edge 1208. In this embodiment, a plurality of teeth 1304 are disposed on leading edge base 1212. These teeth 1304 can be implemented to facilitate guidance of chip carrier 1200 into socket 132. In the embodiment illustrated in FIG. 13, teeth 1304 are implemented as truncated right prisms or pyramidal frustrums. Alternative shapes can be implemented, however, the angled leading edge of teeth 1304 facilitates guidance of carrier 1200. Additionally, in the embodiment illustrated in FIG. 13, teeth 1304 do not extend beyond leading edge base 1212. In this embodiment, teeth 1304 do not extend between contacts 1408 on printed circuit board 1404.

The various embodiments of the invention described above may be implemented using hardware, software or a combination thereof and may be implemented in a computer system or other processing system. In fact, in one embodiment, these elements are implemented using a computer system capable of carrying out the functionality described with respect thereto. An example computer system is such as that illustrated in FIG. 1.

FIG. 16 is a block diagram illustrating a general purpose computer system, including examples of computer readable media for providing computer software or instructions to perform the functionality described herein. The illustrated computer system 1602 includes one or more processors, such as processor 1604. The processor 1604 is connected to a communication bus 1606. Various software embodiments are described in terms of this example computer system. After reading this description, it will become apparent to a person skilled in the relevant art how to implement the invention using other computer systems or computer architectures, including, for example, the architecture illustrated in FIG. 1.

Computer system 1602 also includes a main memory 1608, preferably random access memory (RAM), and can also include a secondary memory 1610. The secondary memory 1610 can include, for example, a hard disk drive 1612 and/or a removable storage drive 1614, representing a floppy disk drive, a magnetic tape drive, an optical disk drive, etc. Removable storage drive 1614 reads from and/or writes to a removable storage medium 1618. Removable storage media 1618, represents a floppy disk, magnetic tape, optical disk, etc. which is read by and written to by removable storage drive 1614. As will be appreciated, the removable storage media 1618 includes a computer-usable storage medium having stored therein computer software and/or data.

In alternative embodiments, secondary memory 1610 may include other similar means for allowing computer programs or other instructions to be loaded into computer system 1602. Such means can include, for example, a removable storage unit 1622 and an interface 1620. Examples of such can include a program cartridge and cartridge interface (such as, for example, that found in video game devices), a removable memory chip (such as, for example, an EPROM, PROM or other memory device) and associated socket, and other removable storage units 1622 and interfaces 1620 which allow software and data to be transferred from the removable storage unit 1622 to computer system 1602. In some embodiments, removable storage unit 1622 may be affixed permanently to removable storage unit interface 1520.

Computer system 1602 can also include a communications interface 1624. Communications interface 1624 allows software and data to be transferred between computer system 1602 and external devices. Examples of communications interface 1624 can include a modem, a network interface (such as an Ethernet card), a communications port, a PCMCIA slot and card, etc. Software and data transferred via communications interface 1624 are in the form of signals which can be electronic, electromagnetic, optical or other signals capable of being received by communications interface 1624. These signals are provided to communications interface via a channel 1628. This channel 1628 carries signals and can be implemented using a wireless medium, wire or cable, fiber optics, or other communications medium. Some examples of a channel can include a phone line, a cellular phone link, an RF link, a network, the Internet, and other communications channels.

In this document, the terms "computer program medium" and "computer usable medium" are used to generally refer to media such as removable storage media 1618, a hard disk installed in hard disk drive 1612, removable storage unit 1622 and signals on channel 1628. These terms can also refer to main memory 1608 where memory 1608 stores a computer program or a part thereof. These computer program products are means for providing software to computer system 1602.

Computer programs or instructions (also called computer control logic) can be stored in main memory 1608 and/or secondary memory 1610. Computer programs can also be received via communications interface 1624. Such computer programs, when executed, enable the computer system 1602 to perform the features of the present invention as discussed herein. In particular, the computer programs, when executed, enable the processor 1604 to perform the features of the present invention. Accordingly, such computer programs represent controllers of the computer system 1602.

In an embodiment where the elements are implemented using software, the software may be stored in a computer program product and loaded into computer system 1602 using removable storage drive 1614, removable storage unit 1622, hard drive 1612 or communications interface 1624. The control logic (software), when executed by the processor 1604, causes the processor 1604 to perform the functions of the invention as described herein.

In another embodiment, the elements are implemented primarily in hardware using, for example, hardware components such as application specific integrated circuits

(ASICs). Implementation of the hardware state machine so as to perform the functions described herein will be apparent to persons skilled in the relevant art(s). Although not a "computer program" in the traditional sense, the hardware components can be thought of as a computer program medium (albeit, perhaps hard-wired) which enables the system to perform the described functions. In yet another embodiment, elements are implemented using a combination of both hardware and software. In this embodiment, the combination of the hardware and software can likewise be thought of as a computer program medium which enables the system to perform the described functions.

While various embodiments of the present invention have been described above, it should be understood that they have been presented by way of example only, and not limitation. Thus, the breadth and scope of the present invention should not be limited by any of the above-described exemplary embodiments, but should be defined only in accordance with the following claims and their equivalents.

WHAT IS CLAIMED IS:

1. A fluorometer for sensing the fluorescence of a sample, comprising:
 - 5 an assay device for carrying the sample to be tested;
 - an optical energy source for exciting the assay device by irradiating the assay device with optical energy;
 - an optical energy detector for detecting the emitted energy from the excited assay device;
 - 10 an encoded label disposed on said assay device to provide encoded sample information relating to the sample carried in the assay device;
 - drive electronics for positioning the assay device to thereby position the assay zones with respect to said optical energy source and said optical energy detector, allowing one of a plurality of assay zones or regions to be tested;
 - 15 a processor for controlling the operation of the test in accordance with test instructions and for processing the emitted energy detected from said region or plurality of regions of the excited sample to determine test results;
 - a ROM chip socket for accepting at least one of a plurality of ROM chips, wherein each said ROM chip stores data relating to one or more assays to be performed; and
 - 20 a communications interface for transferring test information between the fluorometer and an external entity.
2. The fluorometer of claim 1, wherein said ROM chip further stores control and calibration information for calibrating the fluorometer using a control sample.
- 25 3. The fluorometer of claim 1, wherein said sample information includes at least one of the group of a patient identification, a sample identification, a sample type, and an identification of a type of test to be performed on the sample.
- 30 4. The fluorometer of claim 1, wherein said encoded label includes at least one of the group of an optical bar code, a magnetic stripe, an inductive circuit, and alphanumeric characters.

5. The fluorometer of claim 1, wherein said data relating to one or more assays includes at least one of the group of test instructions for a particular type of test, an identification of a particular type of test to be performed, and calibration curves for a test to be performed.

5

6. The fluorometer of claim 1, wherein said test information includes at least one of the group of test results, patient identification, sample identification, and an identification of the test or tests performed on a sample.

10 7. The fluorometer of claim 1, further comprising a power source and wherein said fluorometer is packaged in a hand-held portable case.

8. The fluorometer of claim 1, further comprising a user interface.

15 9. The fluorometer of claim 1, further comprising a storage device for maintaining a record of one or more samples tested and test results for said one or more samples.

20 10. The fluorometer of claim 3, wherein said record comprises at least one of the group of patient identification, a sample identification, a identification of a test performed on said sample, a date and time at which the tests were conducted, and the test results.

25 11. The fluorometer of claim 10, wherein said communications interface communicates said record to a health care facility.

30 12. The fluorometer of claim 1, wherein said optical energy source excites said assay with energy at a first wavelength and said optical energy detector senses energy emitted from said assay device at a second wavelength, wherein said first wavelength is different from said second wavelength.

13. The fluorometer of claim 12, wherein said first wavelength is approximately 670 nanometers and said second wavelength is approximately 760 nanometers.

5 14. The fluorometer of claim 12, further comprising a first filter to cutoff frequencies emitted from said optical energy source which are near said second wavelength, and a second filter to filter frequencies at said first wavelength from being received by said optical energy detector, thereby decreasing the effect of said optical energy source as a source of background for said optical energy detector.

10 15. The fluorometer of claim 14, wherein said first filter cuts off frequencies at wavelengths longer than approximately 690 nanometers and said second filter cuts off frequencies at wavelengths shorter than approximately 710 nanometers.

15 16. A fluorometer for testing a sample drawn from a patient, the fluorometer comprising:

an assay device for performing a test on said sample;

a processor for controlling the operation of the test in accordance with test instructions and for processing data obtained during testing to determine test results; and

20 a removable storage medium for accepting at least one of a plurality of removable storage media, wherein each said removable storage medium includes data sets for one or more assays to be performed.

25 17. The fluorometer according to claim 16, wherein said removable storage medium is a ROM chip.

18. The fluorometer according to claim 16, further comprising an encoded label reader utilized to read an encoded label disposed on an assay device, said label providing encoded sample information relating to the sample disposed in said carrier.

30 19. The fluorometer according to claim 16, further comprising a communications interface for transferring test information between the test device and an external entity.

20. The fluorometer according to claim 16, wherein said fluorometer comprises:

- an optical energy source for exciting the sample to be tested by irradiating the assay device with optical energy; and
- an optical energy detector for detecting the emitted energy from the excited assay device.

21. The fluorometer according to claim 20, wherein said fluorometer further comprises drive electronics for positioning said assay device with respect to said optical energy source and said optical energy detector, allowing at least one of a plurality of sample regions to be tested.

22. The fluorometer of claim 20, wherein said optical energy source excites said assay device with energy at a first wavelength and said optical energy detector senses energy emitted from said assay device at a second wavelength, wherein said first wavelength is different from said second wavelength.

23. The fluorometer of claim 22, wherein said first wavelength is approximately 670 nanometers and said second wavelength is approximately 760 nanometers.

24. The fluorometer of claim 22, further comprising a first filter to cutoff frequencies emitted from said optical energy source which are near said second wavelength, and a second filter to filter frequencies at said first wavelength from being received by said optical energy detector, thereby decreasing the effect of said optical energy source as a source of background for said optical energy detector.

25. The fluorometer of claim 24, wherein said first filter cuts off frequencies at wavelengths longer than approximately 690 nanometers and said second filter cuts off frequencies at wavelengths shorter than approximately 710 nanometers.

26. The fluorometer of claim 16, wherein said first removable storage media storing a test data set for a first type of test can be replaced with a second removable storage media to perform a second type of test.

5 27. The fluorometer of claim 16, further comprising timing means for controlling the execution of the assay, allowing the assay to be conducted without requiring user intervention.

28. A method of performing testing on a sample, comprising the steps of:
10 obtaining a sample to be tested;
programming a test instrument to perform a test on said sample, wherein said programming is accomplished by installing a removable storage medium in said test instrument, wherein said removable storage medium stores data relating to one or more assays to be performed;
15 placing said sample to be tested in an assay device and inserting said assay device into said test instrument; and
executing said test on said assay device according to said programming.

29. The method of claim 28, wherein said sample is blood, serum, plasma,
20 urine, extracted feces, waste water or an extracted ground sample.

30. The method of claim 28, wherein said removable storage medium is a ROM chip.

25 31. The method of claim 28, wherein said step of executing comprises the steps of:
exciting the assay device to be tested by irradiating the assay device with optical energy an optical energy source; and
detecting the emitted energy from the excited assay device using an optical energy
30 detector.

32. The method of claim 28, wherein said step of executing further comprises the step of positioning an assay device with respect to said optical energy source and said

optical energy detector, thereby allowing at least one of a plurality of zones on an assay device to be tested.

33. The method of claim 31, wherein said optical energy source excites said sample with energy at a first wavelength and said optical energy detector senses energy emitted from said sample at a second wavelength, wherein said first wavelength is different from said second wavelength.

34. The method of claim 33, wherein said first wavelength is approximately 670 nanometers and said second wavelength is approximately 760 nanometers.

35. The method of claim 33, further comprising a first filter to cutoff frequencies emitted from said optical energy source which are near said second wavelength, and a second filter to filter frequencies at said first wavelength from being received by said optical energy detector, thereby decreasing the effect of said optical energy source as a source of noise for said optical energy detector.

36. The method of claim 35, wherein said first filter cuts off frequencies at wavelengths longer than approximately 690 nanometers and said second filter cuts off frequencies at wavelengths shorter than approximately 710 nanometers.

37. A computer readable medium, readable by a machine, the medium embodying a program of instructions executable by the machine to perform method steps for testing a sample, the method steps comprising the steps of:

exciting an assay device containing the sample by irradiating the assay device with optical energy from an optical energy source;

detecting the emitted energy from the excited assay device;

positioning the assay device with respect to said optical energy source allowing one or more assay zones of the assay device to be excited and tested.

38. The computer readable medium of claim 37, wherein said method further comprises the steps of:

controlling the operation of the test in accordance with test data; and

processing the emitted energy detected from said region or plurality of regions of the excited sample to determine test results in accordance with test instructions included with said test data.

5 39. The computer readable medium of claim 38, wherein said method further comprises the step of said test data from a ROM chip, wherein said ROM chip includes test instructions for one or more tests.

10 40. The computer readable medium of claim 39, wherein said method further comprises the step of receiving control and calibration information for calibrating the fluorometer using a control sample.

15 41. The computer readable medium of claim 38, wherein said method further comprises the step of reading an encoded label disposed on said assay device to obtain encoded sample information relating to the sample carried in the assay device

20 42. The computer readable medium of claim 38, wherein said method further comprises the step of maintaining a record of one or more samples tested and test results for said one or more samples.

 43. The computer readable medium of claim 42, wherein said method further comprises the step of communicating said record to a health care facility.

25 44. The computer readable medium of claim 38, wherein said optical energy source excites said assay device with energy at a first wavelength and said optical energy detector senses energy emitted from said assay device at a second wavelength, wherein said first wavelength is different from said second wavelength.

30 45. The computer readable medium of claim 44, wherein said first wavelength is approximately 670 nanometers and said second wavelength is approximately 760 nanometers.

46. The computer readable medium of claim 44, further comprising the step of transferring test information between the fluorometer and an external entity via a communications interface.

5 47. The computer readable medium of claim 37, wherein said method further comprises the steps of:

 providing sample information for the sample being tested to a health-care facility at a location remote from said test instrument; and

 receiving test information from a database at said remote health-care facility,
10 wherein said test information is retrieved from said database based on said sample information, and wherein said test information indicates at least a test to be performed for said sample.

 48. A test instrument for in-home testing of patient samples, the test instrument
15 comprising:

 an assay device for accepting a patient sample to be tested;

 means for providing sample information for the patient sample being tested to a health-care facility at a location remote from said test instrument;

 a communications interface for receiving test instructions from a database at said
20 remote health-care facility, wherein said test instructions are retrieved from said database based on said sample information, and wherein said test instructions indicate at least a test to be performed for said patient sample; and

 means for conducting a desired test on said patient sample.

25 49. The test instrument of claim 48, further comprising means for creating a record of said test conducted on said sample.

 50. The test instrument of claim 49, wherein communications interface communicates said record to said health-care facility for subsequent analysis by a health-
30 care professional.

 51. The test instrument of claim 49, further comprising means for storing said record of said test conducted on said sample.

52. The test instrument of claim 48, wherein said sample information includes a patient identification and said test information is retrieved from said database based on said patient identification.

5

53. The test instrument of claim 48, wherein said sample information includes a sample type identification.

54. The test instrument of claim 52, wherein said database includes a flag
10 indicating which one or more tests have been conducted for said patient.

55. The test instrument of claim 48, further comprising an alarm feature for informing the patient when a test is should be conducted.

56. The test instrument of claim 48, wherein said assay device is labeled with an encoded label to provide sample information, and wherein said test instrument further comprises an encoded label reader for reading said sample information for the patient sample from said encoded label.

57. The test instrument of claim 48, wherein said sample information includes information regarding the type of sample being tested, and wherein said test instructions are retrieved based on said sample type.

58. The test instrument of claim 57, wherein said sample information
25 comprises information regarding a type of test to be performed on said sample.

59. The testing instrument of claim 48, wherein instructions for executing said test are downloaded from the remote database with said test information.

60. The testing instrument of claim 48, wherein said test instructions retrieved from said remote database indicate a test to be performed and specific instructions for executing said indicated test are retrieved from local storage.

30

61. The testing instrument of claim 60, wherein said instructions are stored on a removable storage medium.

62. The test instrument of claim 48, wherein said means for conducting said
5 desired test comprises:

an optical energy source for exciting the assay device by irradiating the assay device with optical energy;

an optical energy detector for detecting the emitted energy from the excited assay device;

10 an encoded label disposed on said assay device to provide encoded sample information relating to the sample carried in the assay device;

drive electronics for positioning the assay device to thereby position the assay zones with respect to said optical energy source and said optical energy detector, allowing at least one of a plurality of assay zones or regions to be tested; and

15 a processor for controlling the operation of the test in accordance with test instructions and for processing the emitted energy detected from said region or plurality of regions of the excited sample to determine test results.

63. A test instrument for in-home testing of patient samples, the test instrument
20 comprising:

an assay device for accepting a patient sample to be tested;

means for conducting a designated test on said patient sample;

means for providing results of said designated test to a health-care facility at a location remote from said test instrument.

25

64. The test instrument of claim 63, further comprising means for creating and locally storing a record of said test results.

65. The test instrument of claim 63, further comprising means for providing
30 sample information to said health-care facility.

66. The test instrument of claim 65, further comprising means for receiving test information from a database at said remote health-care facility, wherein said test

information is retrieved from said database based on said sample information, and wherein said test information indicates a test to be performed for said patient sample.

67. The test instrument of claim 65, wherein said sample information
5 comprises an identification of said patient from which said sample was taken.

68. The test instrument of claim 65, wherein said sample information
comprises a sample type identification.

10 69. The test instrument of claim 65, wherein said sample information
comprises an indication of a test to be performed on said patient sample.

70. The test instrument of claim 63, wherein said assay device is labeled with
an encoded label to provide sample information, and wherein said test instrument further
15 comprises means for reading said sample information for the patient sample from said
encoded label.

71. The test instrument of claim 63, wherein instructions for executing said test
are downloaded from the health-care facility based on said sample information.

20 72. The test instrument of claim 63, wherein instructions for executing said test
are retrieved from local storage.

73. The test instrument of claim 63, wherein said means for conducting a
25 designated test comprises:

an optical energy source for exciting the assay device by irradiating the assay
device with optical energy;

an optical energy detector for detecting the emitted energy from the excited assay
device;

30 an encoded label disposed on said assay device to provide encoded sample
information relating to the sample carried in the assay device;

drive electronics for positioning the assay device to thereby position the assay zones with respect to said optical energy source and said optical energy detector, allowing at least one of a plurality of assay zones or regions to be tested; and

5 a processor for controlling the operation of the test in accordance with test instructions and for processing the emitted energy detected from said region or plurality of regions of the excited sample to determine test results.

74. The test instrument of claim 73, further comprising a ROM chip socket for accepting at least one of a plurality of ROM chips, wherein each said ROM chip stores
10 data relating to one or more tests to be performed on said sample.

75. A test instrument for in-home testing of patient samples, the test instrument comprising:

scheduling means indicating a type of test to be performed on a patient sample and
15 a time at which said test should be performed;

means for alerting the patient when said test is scheduled to be performed;

an assay device for accepting a patient sample prepared by the patient;

means for conducting said indicated test on said prepared patient sample; and

a communications interface for providing information pertaining to said test to a
20 health care facility at a remote location.

76. The test instrument of claim 75, wherein said information pertaining to said test comprises an indication of the test conducted and test results.

25 77. The test instrument of claim 75, wherein said scheduling means comprises a scheduling program detailing a scheduled occurrence for performing tests and specifying one or more tests to be performed at each said scheduled occurrence.

78. The test instrument of claim 77, wherein said scheduling program is
30 periodically updated by subsequent downloads via said communications interface.

79. The test instrument of claim 77, wherein said scheduling program retrieves test instructions for a specified test to be performed from local storage or from a remote

database, and provides said test instructions to the test instrument for conducting the specified test.

80. The test instrument of claim 77, wherein said scheduling program is
5 downloaded to the test instrument via said communications interface.

81. The test instrument of claim 78, wherein one or more periodic updates
received via said communications interface adds additional tests to the test schedule,
changes the scheduled occurrence for one or more tests, and/or discontinues one or more
10 currently scheduled tests.

82. The test instrument of claim 77, wherein said scheduling program further
indicates a sample type to be tested at a given scheduled test occurrence.

83. The test instrument of claim 75, wherein once said assay device is prepared
15 with the sample to be tested and the assay device is loaded into the test instrument, the test
is executed and the results reported to the health care facility automatically, without
requiring further patient intervention.

84. An automated testing instrument for testing a sample, said automated
20 testing instrument comprising:

an assay device for accepting a sample to be tested, wherein said assay device is
labeled with an encoded label to provide sample information;

a encoded label reader for reading said sample information for the sample being
25 tested or to be tested;

a communications interface for communicating with a remote database information
regarding a test to be performed or a test which has been performed on said sample.

85. The automated testing instrument of claim 84, wherein said
30 communications interface comprises an interface for retrieving test information from a
remote database, wherein said test information is retrieved based on said sample
information, and wherein said test information indicates at least a test to be performed for
said sample being tested.

86. The automated testing instrument of claim 85, wherein said sample information comprises an identification of a patient from which the sample was drawn, and said patient identification is used to retrieve the test information from said remote database.

5

87. The automated testing instrument of claim 84, further comprising a storage device for maintaining a record of one or more samples tested and test results for said one or more tested samples.

10

88. The automated testing instrument of claim 87, wherein said record comprises at least one of the group of patient identification, a sample identification, a identification of a test performed on said sample, a date and time at which the tests were conducted, and the test results.

15

89. The automated testing instrument of claim 88, wherein said communications interface further communicates said record to a health care facility.

90. The automated testing instrument of claim 84, wherein said sample information includes information regarding the type of sample being tested, and wherein said test information is retrieved based on said sample type.

20

91. The automated testing instrument of claim 84, wherein said test indicated by said retrieved test information is conducted on said sample.

25

92. The automated testing instrument of claim 91, wherein instructions for executing said test are downloaded from the remote database with said test information.

93. The automated testing instrument of claim 91, wherein instructions for executing said test are retrieved from local storage based on said test information.

30

94. A method for performing automated testing of samples, the method comprising the steps of:

47

accepting an assay device including a patient sample to be tested;
reading an encoded label on said assay device, said encoded label indicating the
patient from which the sample was drawn;

determining which one or more of a plurality of tests to perform on said sample
5 based on said encoded label; and
automatically conducting said one or more tests on the patient sample.

95. The method of claim 94, wherein said encoded label further indicates the
type of sample being tested.

10

96. The method of claim 94, wherein said encoded label further indicates a type
of test for said patient sample.

97. The method of claim 94, further comprising the step of recording a test
15 record indicating at least one of the type of test conducted, the identification of the patient
and results of the conducted test.

98. The method of claim 97, further comprising the step of forwarding said test
record to a health care facility via a communications interface.

20

99. The method of claim 98, further comprising the step of storing information
from said test record in a health care database at said health care facility.

100. The method of claim 94, wherein said step of determining which one or
25 more of a plurality of tests to perform comprises the step of using said patient indication to
look up a test identification in a database.

101. The method of claim 96, wherein said sample type is at least one of the
group of blood or urine.

30

102. The method of claim 100, wherein said database is a local database.

103. The method of claim 100, wherein said database is a remote database at a health care facility.

104. The method of claim 94, further comprising the step of receiving a scheduling program detailing a scheduled occurrence for performing one or more tests and specifying one or more tests to be performed at each said scheduled occurrence.

105. The method of claim 100, further comprising the step of periodically updating said scheduling program by subsequent downloads via said communications interface or via a removable storage medium.

106. The method of claim 100, further comprising the step of retrieving test instructions for a specified test to be performed from local storage or from a remote database, and provides said test instructions to the test instrument for conducting the specified test.

107. The method of claim 100, further comprising the step of downloading said scheduling program to the test instrument via a communications interface.

108. The method of claim 100, further comprising the step of receiving periodic updates to said scheduling program, wherein said periodic updates add additional tests to the test schedule, change the scheduled occurrence for one or more tests, and/or discontinue one or more currently scheduled tests.

109. The method of claim 97, wherein once said assay device is loaded into the test instrument, the test type is determined, the test executed and the results recorded automatically, without requiring further patient intervention.

110. The method of claim 94, further comprising the step of forwarding information regarding a conducted test to a health care facility at a remote location.

111. The method of claim 110, wherein said patient is identified by an ANI.

112. A computer readable medium, embodying a program of instructions for causing a machine to perform a method for performing automated testing of samples, the method comprising the steps of:

- accepting an assay device including a patient sample to be tested;
- 5 reading an encoded label on said assay device, said encoded label indicating sample information regarding said patient sample;
- determining which one or more of a plurality of tests to perform on said sample based on said encoded label; and
- automatically conducting said one or more tests on the patient sample.

10

113. The computer readable medium of claim 112, wherein said step of determining comprises the steps of

- providing sample information for the sample being tested to a health-care facility at a location remote from said test instrument; and
- 15 receiving test information from a database at said remote health-care facility, wherein said test information is retrieved from said database based on said sample information, and wherein said test information indicates at least a test to be performed for said sample.

20

114. The computer readable medium of claim 112, wherein said sample information indicates the type of sample being tested.

25

115. The computer readable medium of claim 112, wherein said sample information indicates an identification of the patient from which said patient sample was drawn.

30

116. The computer readable medium of claim 112, wherein said sample information indicates a type of test which is to be performed or which has been performed on said sample.

117. The computer readable medium of claim 113, wherein said patient is identified based on an ANI.

118. The computer readable medium of claim 112, further comprising the step of recording a test record indicating at least one of the type of test conducted, the identification of the patient and results of the conducted test.

5 119. The computer readable medium of claim 118, further comprising the step of forwarding said test record to a health care facility via a communications interface.

120. The computer readable medium of claim 119, further comprising the step of storing information from said test record in a health care database at said health care
10 facility.

121. The computer readable medium of claim 112, wherein said step of determining which one or more of a plurality of tests to perform comprises the step of using said sample information to look up a test identification in a database.

15 122. The computer readable medium of claim 121, wherein said database is a local database.

123. The computer readable medium of claim 121, wherein said database is a
20 remote database at a health care facility.

124. The computer readable medium of claim 112, further comprising the step of receiving a scheduling program detailing a scheduled occurrence for performing one or more tests and specifying one or more tests to be performed at each said scheduled
25 occurrence.

125. The computer readable medium of claim 124, further comprising the step of periodically updating said scheduling program by subsequent downloads via said communications interface or via a removable storage medium.

30 126. The computer readable medium of claim 124, further comprising the step of retrieving test instructions for a specified test to be performed from local storage or

from a remote database, and provides said test instructions to the test instrument for conducting the specified test.

127. The computer readable medium of claim 124, further comprising the step
5 of downloading said scheduling program to the test instrument via a communications interface.

128. The computer readable medium of claim 124, further comprising the step
10 of receiving periodic updates to said scheduling program, wherein said periodic updates add additional tests to the test schedule, change the scheduled occurrence for one or more tests, and/or discontinue one or more currently scheduled tests.

129. The computer readable medium of claim 112, wherein once said assay
15 device is loaded into the test instrument, the test type is determined, the test executed and the results recorded automatically, without requiring further patient intervention.

130. A fluorometer for sensing the fluorescence of a sample, comprising:
an assay device for carrying the sample to be tested;
an optical energy source for exciting the assay device by irradiating the assay
20 device with optical energy;
an optical energy detector for detecting the emitted energy from the excited assay device;
drive electronics for positioning the assay device to thereby position the assay zones with respect to said optical energy source and said optical energy detector, allowing
25 at least one of a plurality of assay zones or regions to be tested; and
a processor for controlling the operation of the test in accordance with test instructions and for processing the emitted energy detected from said region or plurality of regions of the excited sample to determine test results.

30 131. The fluorometer of claim 130, further comprising a ROM chip socket for accepting at least one of a plurality of ROM chips which store data relating to one or more assays to be performed, and wherein said ROM chips further store control and calibration information for calibrating the fluorometer using a control sample.

132. The fluorometer of claim 131, wherein said sample information includes at least one of the group of a patient identification, a sample identification, a sample type, and an identification of a type of test to be performed on the sample.

5

133. The fluorometer of claim 131, further comprising an encoded label to provide encoded sample information, wherein said encoded label includes at least one of the group of an optical bar code, a magnetic stripe, an inductive circuit, and alphanumeric characters.

10

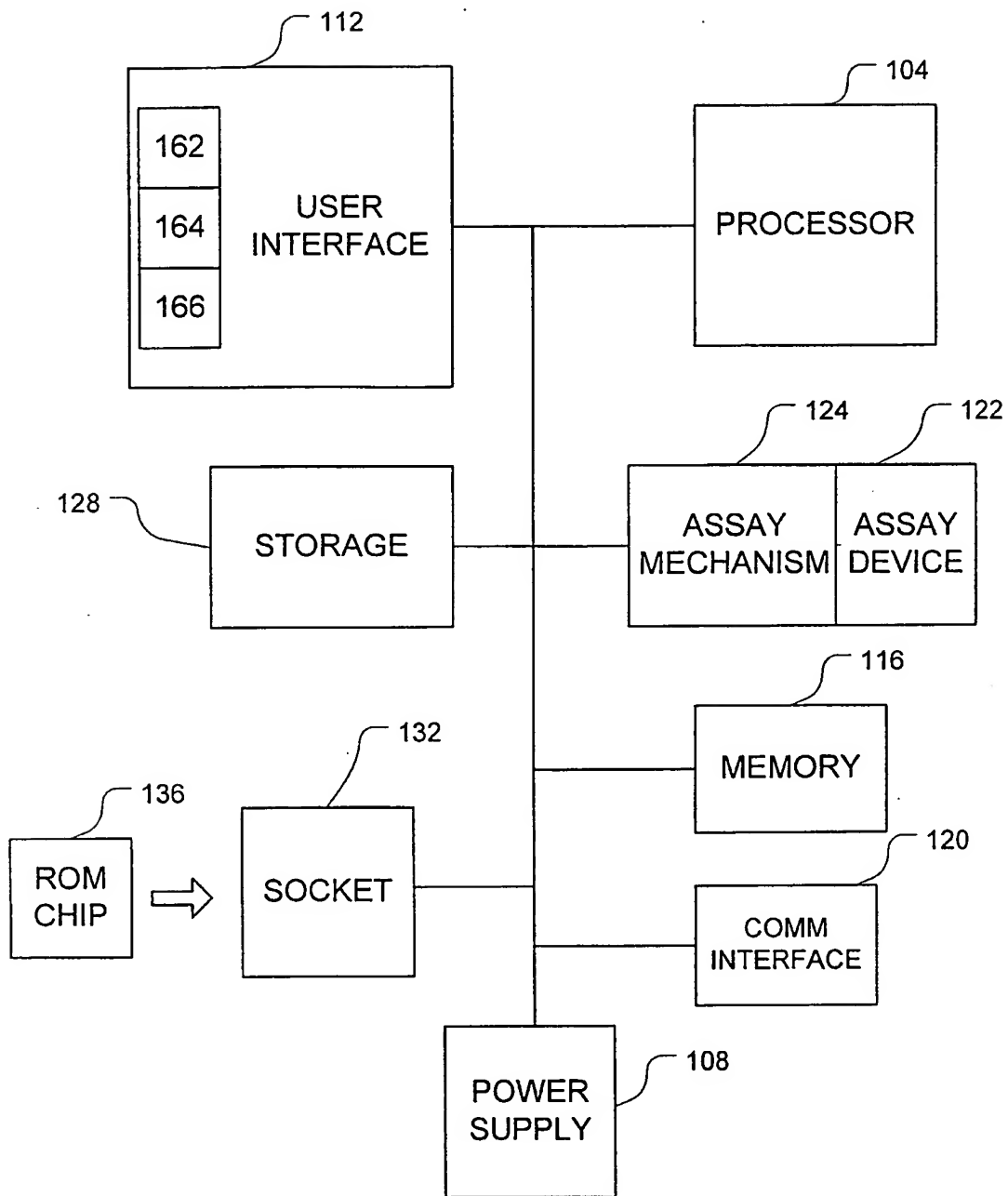
134. The fluorometer of claim 131, further comprising:

a processor for controlling the operation of the test in accordance with test instructions and for processing the emitted energy detected from said region or plurality of regions of the excited sample to determine test results; and

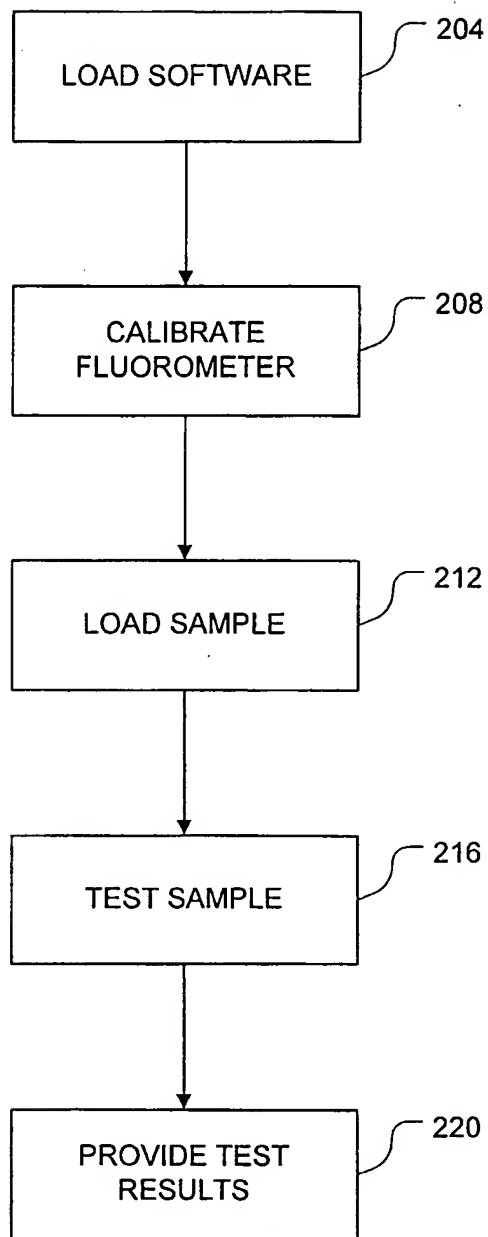
15

a communications interface for transferring test information between the fluorometer and an external entity.

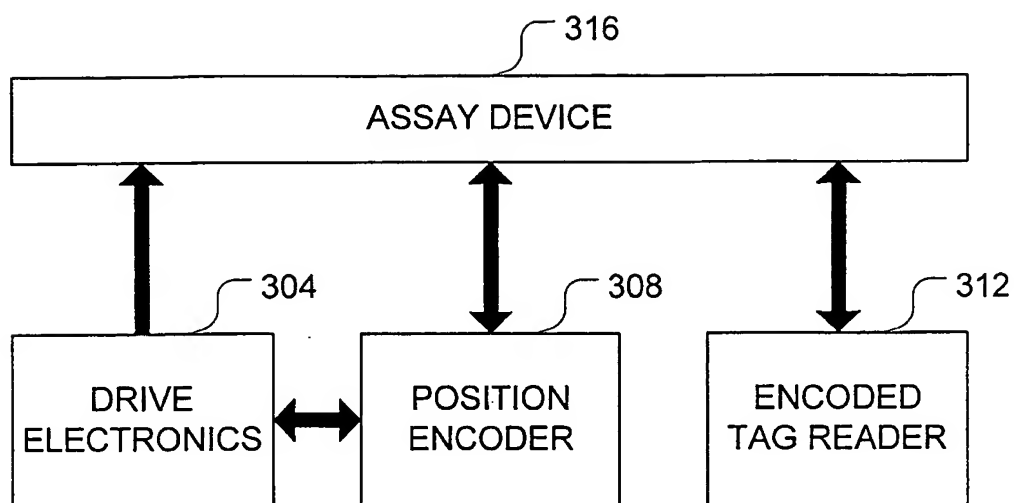
135. The fluorometer of claim 134, wherein said communications interface communicates said test information to a health care facility.

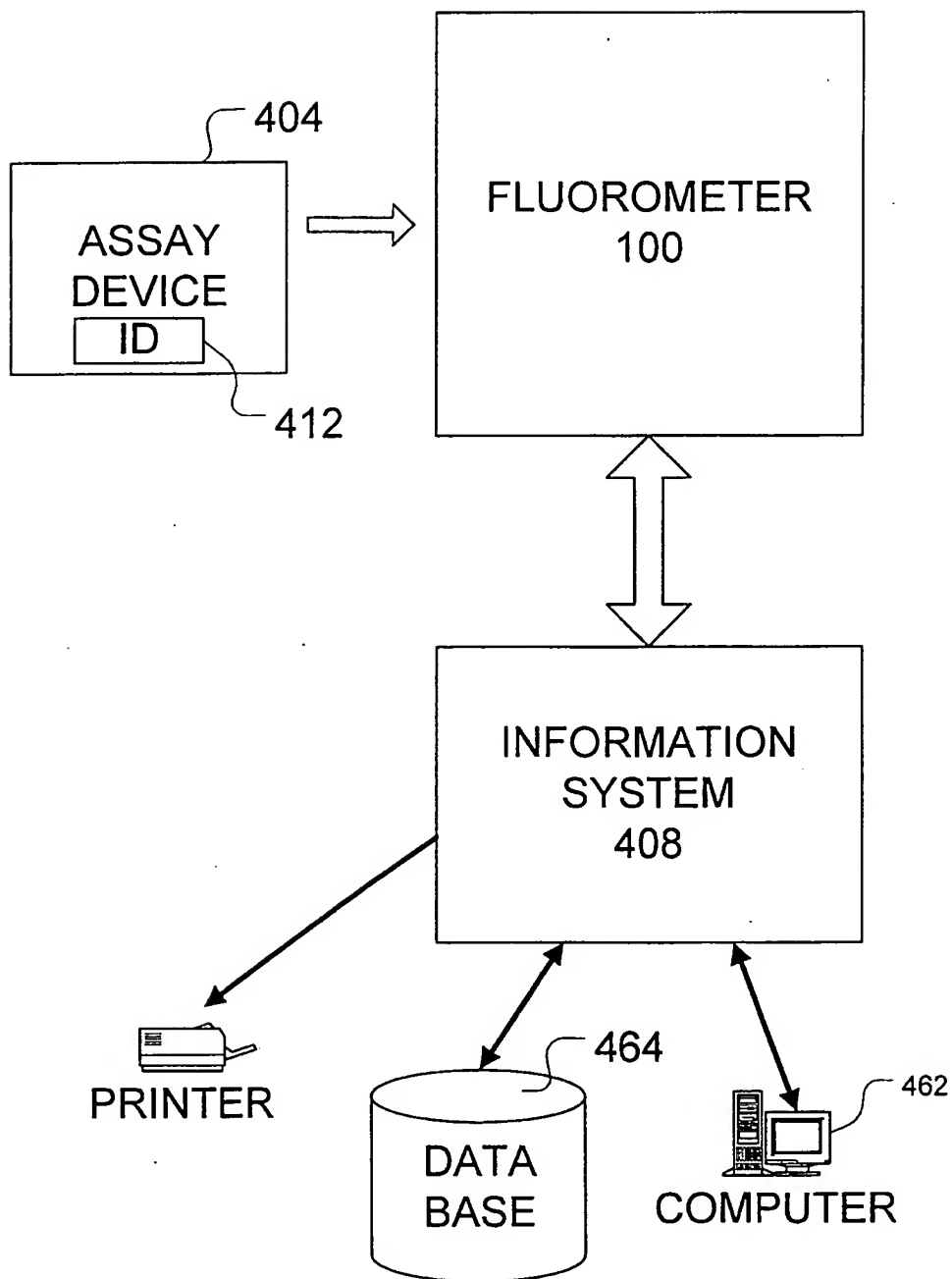
*FIG. 1*

2/13

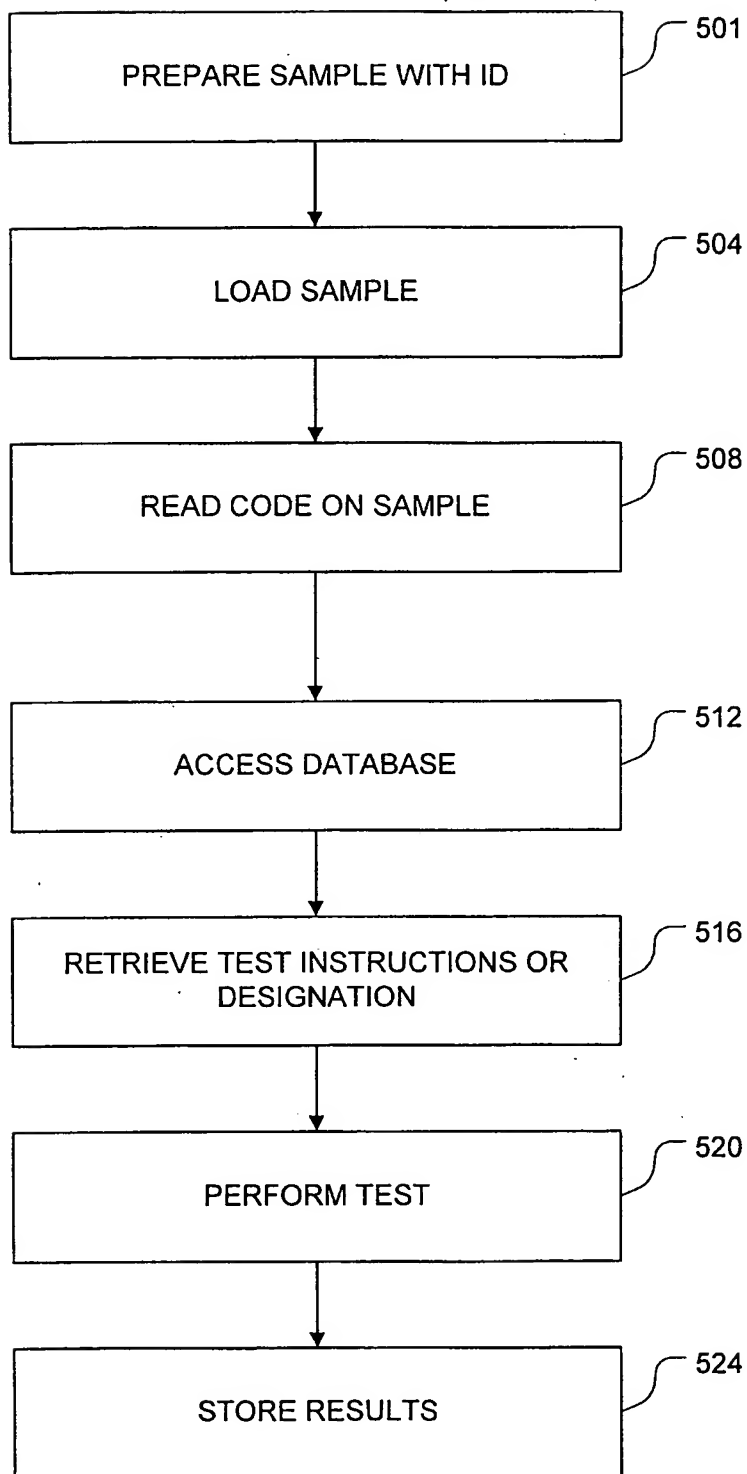
*FIG. 2*

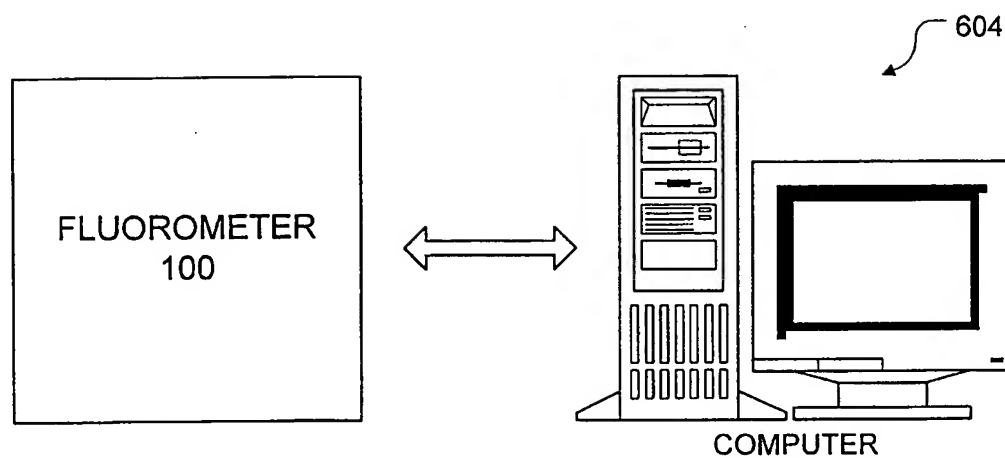
3/13

*FIG. 3*

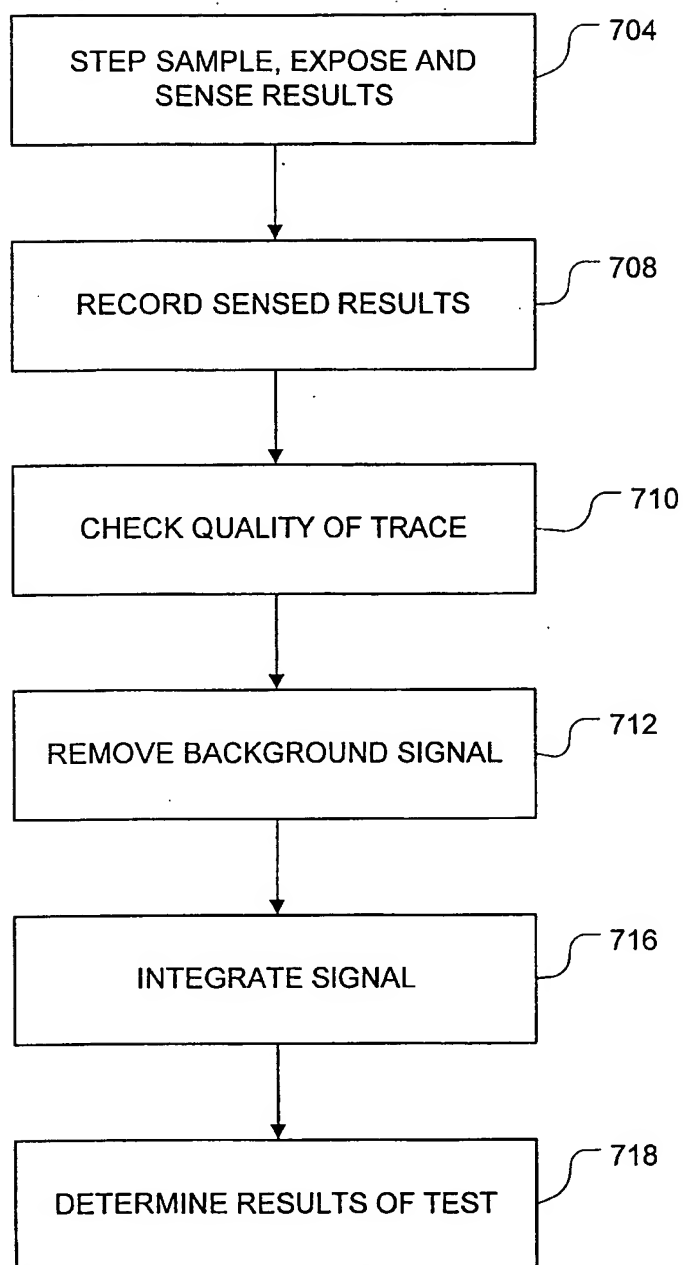
*FIG. 4*

5/13

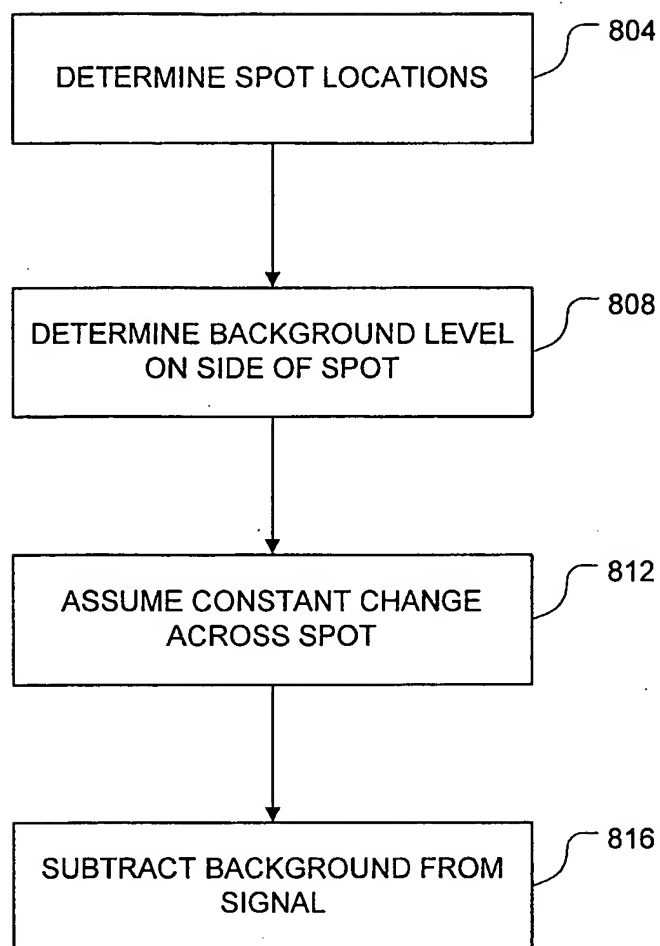
*FIG. 5*

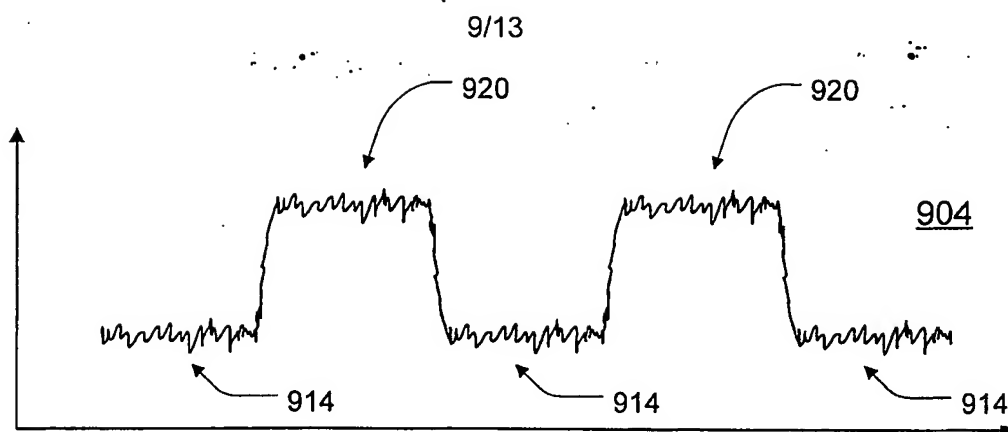
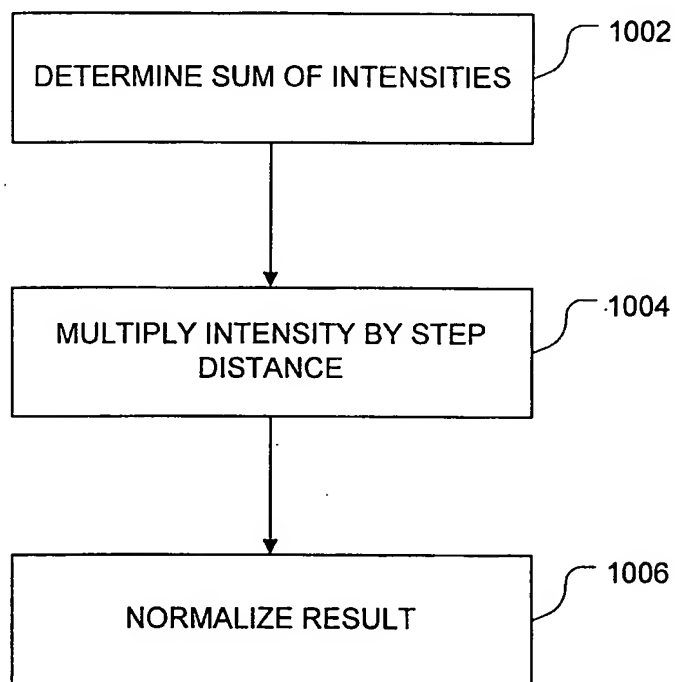
*FIG. 6*

7/13

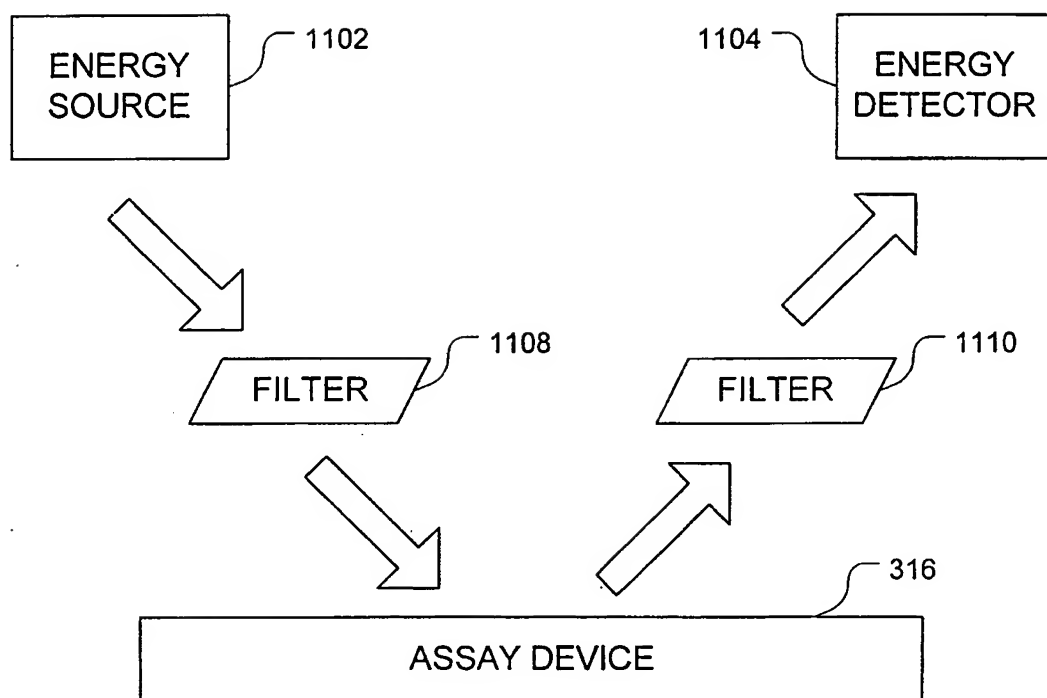
*FIG. 7*

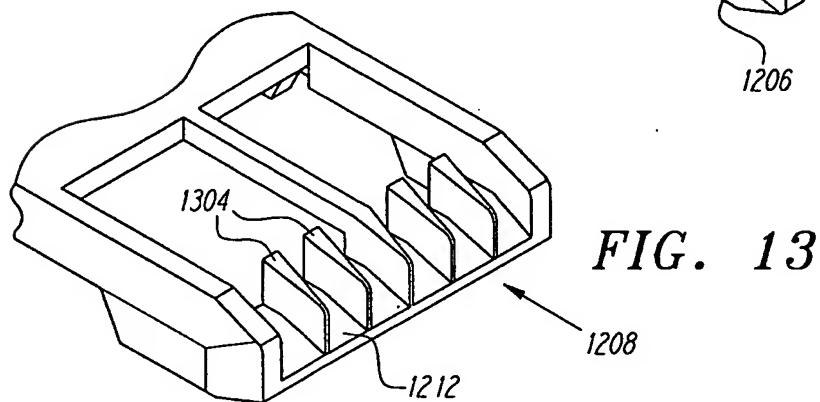
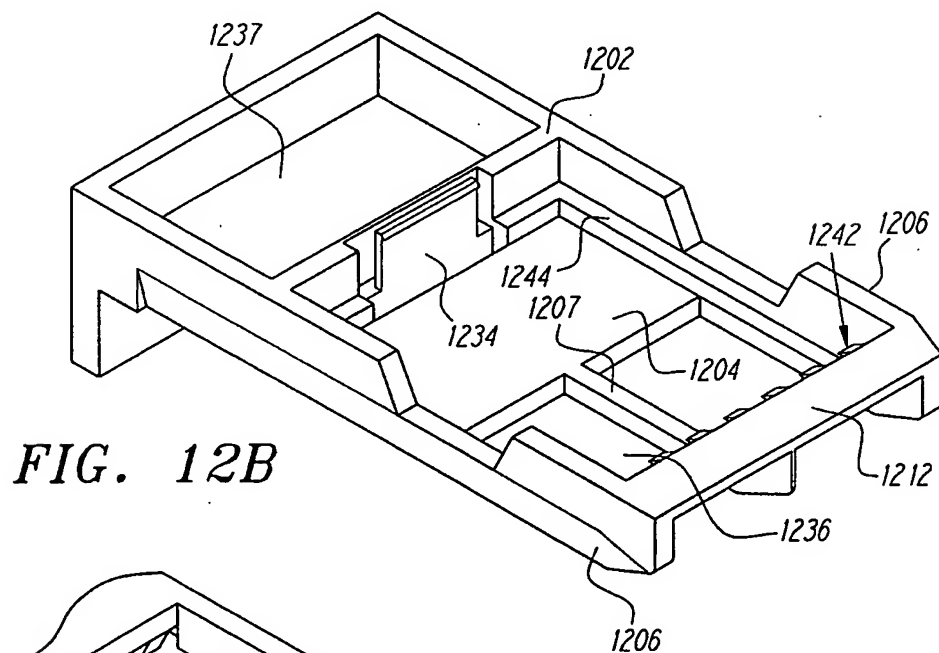
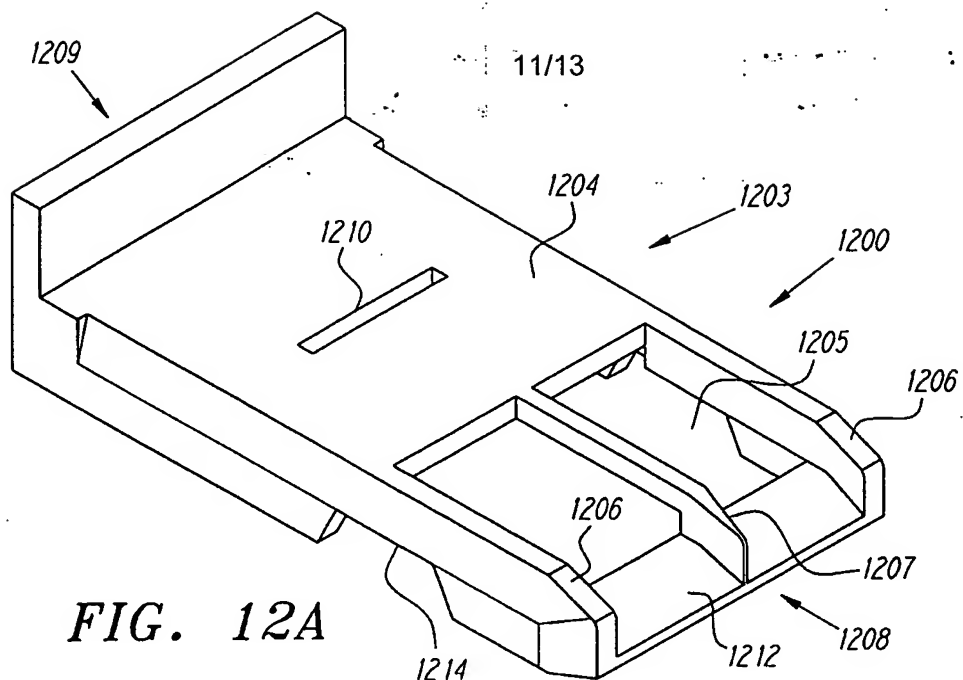
8/13

*FIG. 8*

*FIG. 9**FIG. 10*

10/13

*FIG. 11*



12/13

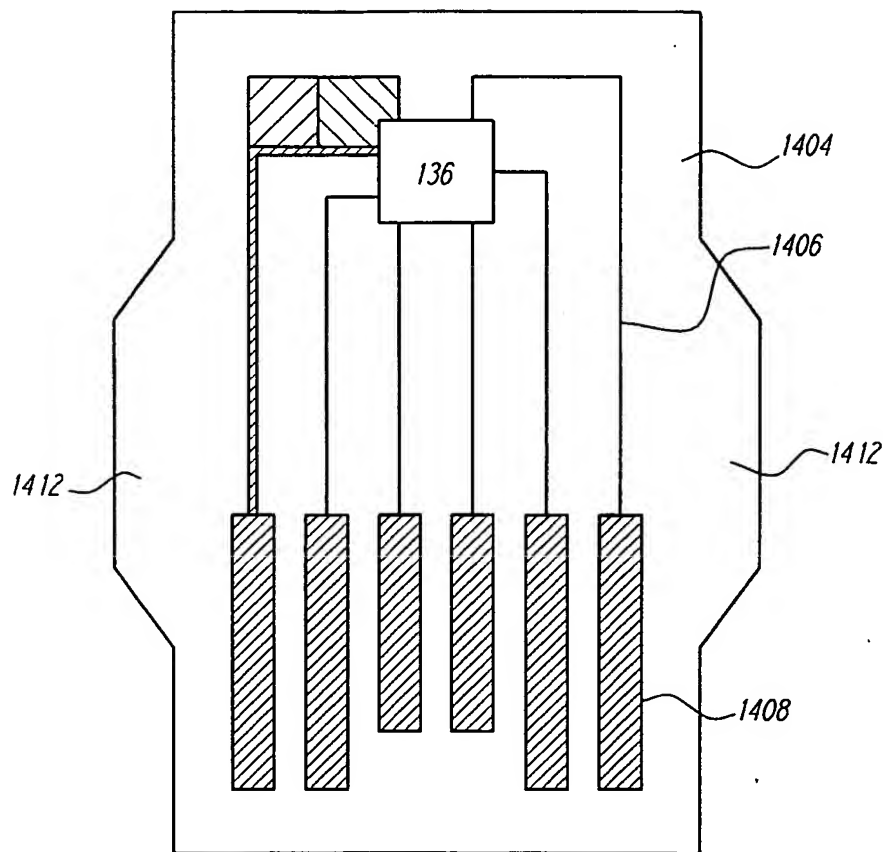


FIG. 14

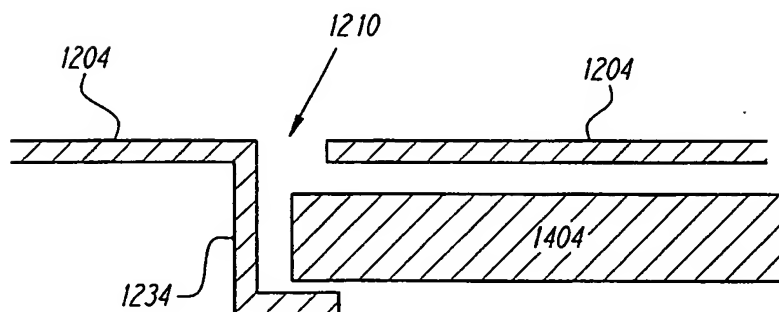
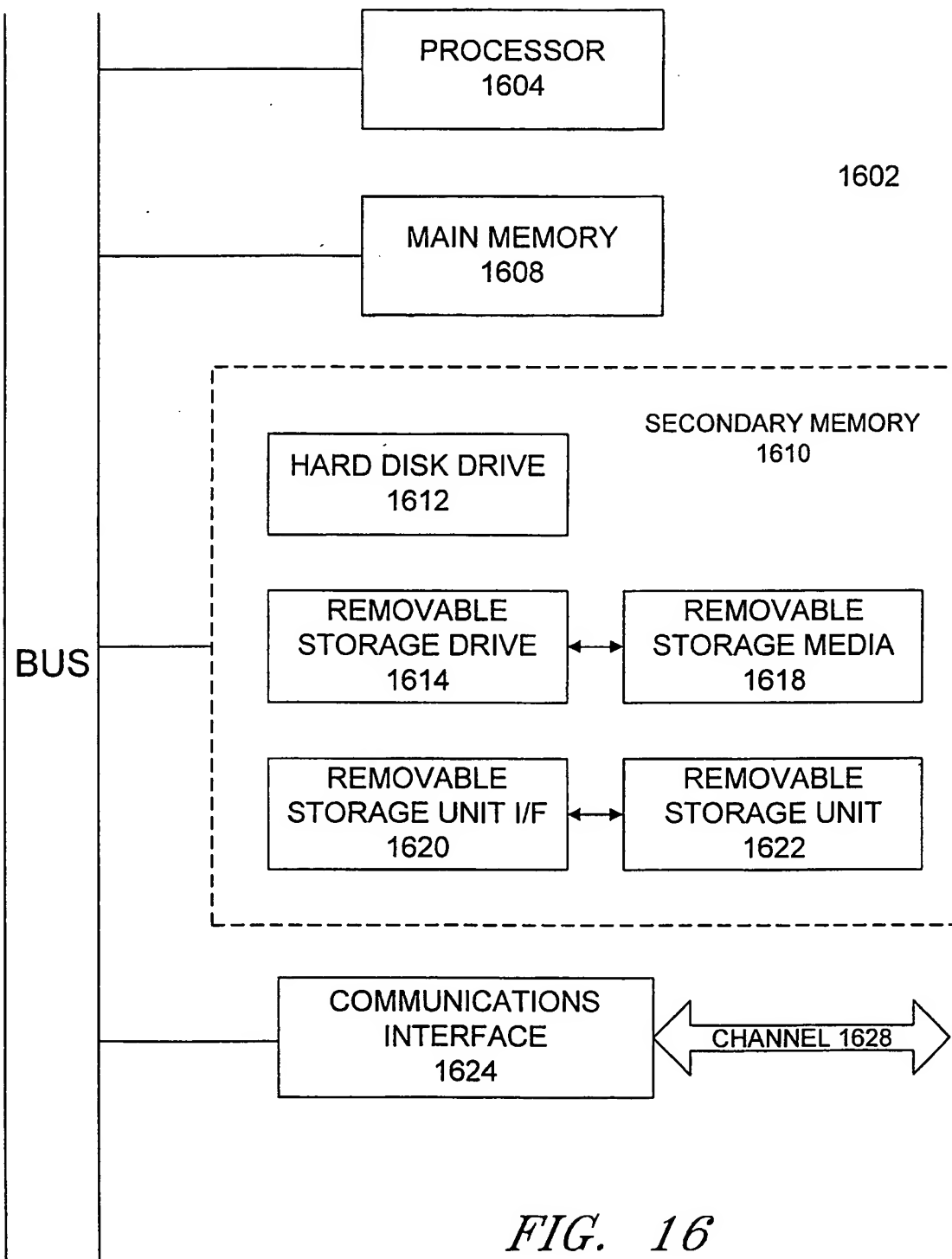


FIG. 15

13/13



INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 98/26576

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 G01N21/64

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 G01N A61B G06F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 5 695 949 A (GALEN ROBERT S ET AL) 9 December 1997 see column 2, line 31 - line 40 see column 11, line 53 - line 54 see column 12, line 40 - line 66; figure 4 see column 13, line 15 - line 59; figures 2-4	1-135
Y	US 5 594 906 A (ANDERSON JOE E ET AL) 14 January 1997 see column 1, line 6 - line 23 see column 3, line 33 - line 35 see column 8, line 24 - line 39	1-135
	--- -/-- ---	



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"A" document member of the same patent family

Date of the actual completion of the international search

29 April 1999

Date of mailing of the international search report

12/05/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Hocquet, A

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 98/26576

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 3 907 503 A (BETTS CLEO ELMER ET AL) 23 September 1975 see column 2, line 65 - column 3, line 41 see column 5, line 24 - line 59; figure 1 see column 6, line 19 - line 54 ----	1
A	US 5 597 532 A (CONNOLLY JAMES) 28 January 1997 see column 2, line 20 - line 67; figure 1 see column 5, line 30 - line 32 see column 10, line 28 - line 36 ----	1
X	US 5 053 199 A (JACKSON CHARLES A ET AL) 1 October 1991 see column 1, line 26 - column 2, line 21 see column 2, line 54 - column 3, line 6 ----	28-47, 112-129
A	US 5 281 395 A (MARKART ERNST ET AL) 25 January 1994 see column 1, line 15 - line 28 see column 3, line 5 - line 66 ----	1-135 28-47, 112-129
A	WO 96 29367 A (BIOSITE DIAGNOSTICS INC) 26 September 1996 cited in the application see page 2, line 16 - page 3, line 13 see page 26, line 1 - line 14 see page 4, line 7 - line 12 see page 10, line 12 - line 15 ----	1-135
A	US 5 030 009 A (ANDO ITSURO ET AL) 9 July 1991 see column 1, line 35 - line 57 see column 5, line 33 - line 68; figure 1 see column 6, line 44 - line 68; figure 2 ----	1-135
A	US 5 458 852 A (BUECHLER KENNETH F) 17 October 1995 cited in the application see column 11, line 29 - line 31 see column 12, line 52 - column 13, line 2; figure 3 ----	1
X	WO 97 28737 A (NOKIA MOBILE PHONES LTD ;HEINONEN PEKKA (FI); OKKONEN HARRI (FI)) 14 August 1997 see page 4, line 29 - page 5, line 30; figures see page 10, line 19 - line 23 ----	48-111
X	EP 0 483 595 A (MILES INC) 6 May 1992 see column 7, line 24 - column 8, line 15; figure 1 -----	48-111

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 98/26576

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5695949 A	09-12-1997	CA 2215914 A EP 0819180 A JP 11503025 T WO 9631619 A	10-10-1996 21-01-1998 23-03-1999 10-10-1996
US 5594906 A	14-01-1997	US 5371687 A CA 2147484 A DE 69323094 D EP 0680645 A EP 0836135 A JP 8504045 T WO 9412950 A	06-12-1994 09-06-1994 25-02-1999 08-11-1995 15-04-1998 30-04-1996 09-06-1994
US 3907503 A	23-09-1975	AR 213268 A AT 357350 B AT 40775 A AU 7596274 A BE 824370 A BR 7500327 A CA 1033651 A CA 1047379 A CH 588697 A CS 209419 B DD 116678 A DE 2502013 A DK 687774 A FI 750055 A,B, FR 2258630 A GB 1461319 A IE 40749 B JP 1043118 C JP 50104991 A JP 55033544 B JP 56000656 A NL 7417004 A SE 7500331 A YU 13775 A ZA 7407835 A	15-01-1979 10-07-1980 15-11-1979 03-06-1976 02-05-1975 04-11-1975 27-06-1978 30-01-1979 15-06-1977 31-12-1981 05-12-1975 11-09-1975 13-10-1975 22-07-1975 18-08-1975 13-01-1977 15-08-1979 23-04-1981 19-08-1975 01-09-1980 07-01-1981 23-07-1975 22-07-1975 27-04-1983 28-04-1976
US 5597532 A	28-01-1997	EP 0750739 A WO 9613707 A	02-01-1997 09-05-1996
US 5053199 A	01-10-1991	AT 152844 T AU 5189690 A CA 2010400 A,C DE 69030657 D DE 69030657 T EP 0460074 A ES 2100881 T JP 4506721 T WO 9010236 A	15-05-1997 26-09-1990 21-08-1990 12-06-1997 11-09-1997 11-12-1991 01-07-1997 19-11-1992 07-09-1990
US 5281395 A	25-01-1994	DE 4041905 A AT 119292 T DE 59104792 D EP 0492326 A ES 2071198 T GR 3015727 T	02-07-1992 15-03-1995 06-04-1995 01-07-1992 16-06-1995 31-07-1995

INTERNATIONAL SEARCH REPORT

information on patent family members

Inter national Application No

PCT/US 98/26576

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5281395 A		JP 7005109 A	10-01-1995
		JP 7035998 B	19-04-1995
WO 9629367 A	26-09-1996	AU 5318896 A	08-10-1996
		CA 2215727 A	26-09-1996
		EP 0820489 A	28-01-1998
		JP 10508897 T	02-09-1998
		US 5824799 A	20-10-1998
US 5030009 A	09-07-1991	JP 2077302 C	09-08-1996
		JP 2243944 A	28-09-1990
		JP 7119690 B	20-12-1995
US 5458852 A	17-10-1995	AU 4596593 A	30-12-1993
		EP 0596104 A	11-05-1994
		JP 6509424 T	20-10-1994
		WO 9324231 A	09-12-1993
		US 5885527 A	23-03-1999
WO 9728737 A	14-08-1997	FI 960637 A	13-08-1997
		AU 1726797 A	28-08-1997
		EP 0883371 A	16-12-1998
		US 5840020 A	24-11-1998
EP 0483595 A	06-05-1992	US 5251126 A	05-10-1993
		AU 637711 B	03-06-1993
		AU 8020591 A	07-05-1992
		CA 2051914 A,C	30-04-1992
		JP 4348747 A	03-12-1992

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☐ BLACK BORDERS
- ☐ IMAGE CUT OFF AT TOP, BOTTOM OR SIDES
- ☐ FADED TEXT OR DRAWING
- ☐ BLURRED OR ILLEGIBLE TEXT OR DRAWING
- ☐ SKEWED/SLANTED IMAGES
- ☐ COLOR OR BLACK AND WHITE PHOTOGRAPHS
- ☐ GRAY SCALE DOCUMENTS
- ☒ LINES OR MARKS ON ORIGINAL DOCUMENT
- ☐ REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY
- ☐ OTHER: _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.